10/62,3171 Thomas McKenzie

Connecting via Winsock to STN

Welcome to STN International! Enter x:x

LOGINID:ssspta1611txm

PASSWORD:
TERMINAL (ENTER 1, 2, 3, OR ?):2

* * * * * * * * * * Welcome to STN International * * * * * * * * * *

NEWS	1		Web Page URLs for STN Seminar Schedule - N. America
NEWS	2		"Ask CAS" for self-help around the clock
NEWS	3	JAN 27	Source of Registration (SR) information in REGISTRY updated and searchable
NEWS	4	JAN 27	A new search aid, the Company Name Thesaurus, available in CA/CAplus
NEWS	5	FEB 05	German (DE) application and patent publication number format changes
NEWS	6	MAR 03	MEDLINE and LMEDLINE reloaded
NEWS	7	MAR 03	MEDLINE file segment of TOXCENTER reloaded
NEWS	8	MAR 03	FRANCEPAT now available on STN
NEWS	9	MAR 29	Pharmaceutical Substances (PS) now available on STN
NEWS	10	MAR 29	WPIFV now available on STN
NEWS	11	MAR 29	New monthly current-awareness alert (SDI) frequency in RAPRA
NEWS	12	APR 26	PROMT: New display field available
NEWS	13	APR 26	IFIPAT/IFIUDB/IFICDB: New super search and display field available
NEWS	14	APR 26	LITALERT now available on STN
NEWS	15	APR 27	NLDB: New search and display fields available
NEWS	16	May 10	PROUSDDR now available on STN
NEWS	17	May 10	PROUSDDR: One FREE connect hour, per account, in both May and June 2004

NEWS EXPRESS	MARCH 31 CURRENT WINDOWS VERSION IS V7.00A, CURRENT MACINTOSH VERSION IS V6.0c(ENG) AND V6.0Jc(JP), AND CURRENT DISCOVER FILE IS DATED 26 APRIL 2004
NEWS HOURS	STN Operating Hours Plus Help Desk Availability
NEWS INTER	General Internet Information
NEWS LOGIN	Welcome Banner and News Items
NEWS PHONE	Direct Dial and Telecommunication Network Access to STN
NEWS WWW	CAS World Wide Web Site (general information)

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* * * * * * * * * * * * * * STN Columbus * * * * * * * * * * * * * * * *

FILE 'HOME' ENTERED AT 14:20:01 ON 11 MAY 2004

=> file reg
FILE 'REGISTRY' ENTERED AT 14:21:06 ON 11 MAY 2004
USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT.
PLEASE SEE "HELP USAGETERMS" FOR DETAILS.

Property values tagged with IC are from the ZIC/VINITI data file provided by InfoChem.

STRUCTURE FILE UPDATES: 9 MAY 2004 HIGHEST RN 680971-82-8
DICTIONARY FILE UPDATES: 9 MAY 2004 HIGHEST RN 680971-82-8

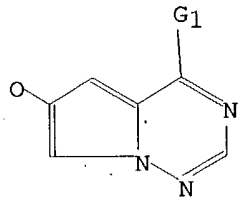
TSCA INFORMATION NOW CURRENT THROUGH JANUARY 6, 2004

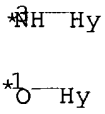
Please note that search-term pricing does apply when conducting SmartSELECT searches.

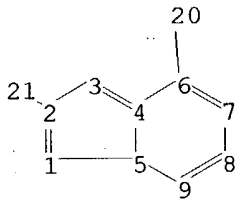
Crossover limits have been increased. See HELP CROSSOVER for details.

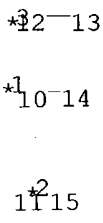
Experimental and calculated property data are now available. For more information enter HELP PROP at an arrow prompt in the file or refer to the file summary sheet on the web at:
http://www.cas.org/ONLINE/DBSS/registryss.html

=>
Uploading C:\Program Files\Stnexp\Queries\10623171.str









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10 11 12 13 14 15 20 21
ring nodes :
1 2 3 4 5 6 7 8 9
chain bonds :
2-21 6-20 10-14 11-15 12-13
ring bonds :
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exact/norm bonds :
1-2 1-5 2-3 2-21 3-4 4-5 4-6 5-9 6-7 6-20 7-8 8-9 10-14 11-15 12-13

G1:OH,Cl,[*1],[*2],[*3]

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Match level :
1:Atom  2:Atom  3:Atom  4:Atom  5:Atom  6:Atom  7:Atom  8:Atom  9:Atom  10:CLASS
11:CLASS  12:CLASS  13:Atom  14:Atom  15:Atom  20:CLASS  21:CLASS
Generic attributes :
13:
Saturation              : Unsaturated
Number of Carbon Atoms : 7 or more
Number of Hetero Atoms : less than 2
Type of Ring System     : Polycyclic
14:
Saturation              : Unsaturated
Number of Carbon Atoms : 7 or more
Number of Hetero Atoms : less than 2
Type of Ring System     : Polycyclic
15:
Saturation              : Unsaturated
Number of Carbon Atoms : 7 or more
Number of Hetero Atoms : less than 2
Type of Ring System     : Polycyclic



L1      STRUCTURE UPLOADED

=> s l1
SAMPLE SEARCH INITIATED 14:21:39 FILE 'REGISTRY'
SAMPLE SCREEN SEARCH COMPLETED -    115 TO ITERATE

100.0% PROCESSED     115 ITERATIONS                          10 ANSWERS
SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS:  ONLINE  **COMPLETE**
                        BATCH   **COMPLETE**
PROJECTED ITERATIONS:         1657 TO     2943
PROJECTED ANSWERS:              11 TO      389

L2           10 SEA SSS SAM L1

=> d scan

L2   10 ANSWERS   REGISTRY  COPYRIGHT 2004 ACS on STN
IN   Pyrrolo[2,1-f][1,2,4]triazine-6-propanoic acid, 4-chloro-5-methyl-, methyl
     ester (9CI)
MF   C11 H12 Cl N3 O2
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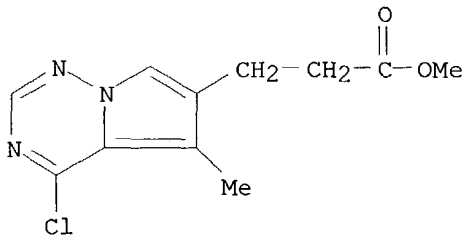
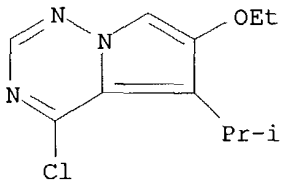


PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

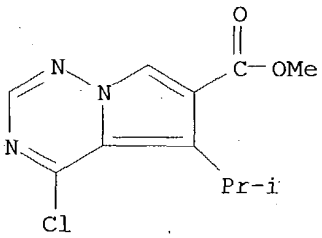
HOW MANY MORE ANSWERS DO YOU WISH TO SCAN? (1):2

L2 10 ANSWERS REGISTRY COPYRIGHT 2004 ACS on STN
IN Pyrrolo[2,1-f][1,2,4]triazine, 4-chloro-6-ethoxy-5-(1-methylethyl)- (9CI)
MF C11 H14 Cl N3 O



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L2 10 ANSWERS REGISTRY COPYRIGHT 2004 ACS on STN
IN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 4-chloro-5-(1-methylethyl)-, methyl ester (9CI)
MF C11 H12 Cl N3 O2



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

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HOW MANY MORE ANSWERS DO YOU WISH TO SCAN? (1):0

=> s l1 full
FULL SEARCH INITIATED 14:23:56 FILE 'REGISTRY'
FULL SCREEN SEARCH COMPLETED -   2229 TO ITERATE

100.0% PROCESSED    2229 ITERATIONS                          186 ANSWERS
SEARCH TIME: 00.00.01

L3          186 SEA SSS FUL L1

=> file caplus
FILE 'CAPLUS' ENTERED AT 14:24:05 ON 11 MAY 2004
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FILE COVERS 1907 - 11 May 2004  VOL 140 ISS 20
FILE LAST UPDATED: 10 May 2004  (20040510/ED)

  This file contains CAS Registry Numbers for easy and accurate
  substance identification.


=> s wo-2000071129?/pn
L4            1 WO-2000071129?/PN
                (WO2000071129/PN)

=> s wo-2004009784?/pn
L5            1 WO-2004009784?/PN
                (WO2004009784/PN)

=> s l3 and l4
             17 L3
L6            1 L3 AND L4

=> d cbib pi hitstr

L6   ANSWER 1 OF 1  CAPLUS  COPYRIGHT 2004 ACS on STN
2000:841986  Document No. 134:17506  Preparation of pyrrolotriazines as
     kinases inhibitors for treating inflammation, cancer, and proliferative
     diseases.  Hunt, John T.; Bhide, Rajeev S.; Borzilleri, Robert M.; Qian,
     Ligang (Bristol-Myers Squibb Company, USA).  PCT Int. Appl. WO 2000071129
     A1 20001130, 130 pp.  DESIGNATED STATES: W:  AE, AL, AM, AT, AU, AZ, BA,
     BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD,
     GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR,
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LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM; RW: AT, BE, BF, BJ, CF, CG, CH, CI, CM, CY, DE, DK, ES, FI, FR, GA, GB, GR, IE, IT, LU, MC, ML, MR, NE, NL, PT, SE, SN, TD, TG. (English). CODEN: PIXXD2. APPLICATION: WO 2000-US13420 20000516. PRIORITY: US 1999-PV135265 19990521; US 2000-PV193727 20000331.

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE	
PI	WO 2000071129	A1	20001130	WO 2000-US13420	20000516	<--
	W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM					
	RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG					
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	JP 2003500359	T2	20030107	JP 2000-619433	20000516	
	NO 2001005650	A	20011120	NO 2001-5650	20011120	
	ZA 2001009577	A	20030220	ZA 2001-9577	20011120	

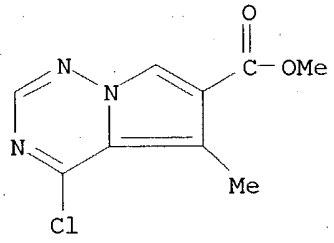
IT **310442-40-1P 310442-94-5P**

RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); RCT (Reactant); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); RACT (Reactant or reagent); USES (Uses)

(preparation of pyrrolotriazines as kinases inhibitors useful in treating inflammation, cancer, and proliferative diseases)

RN 310442-40-1 CAPLUS

CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 4-chloro-5-methyl-, methyl ester (9CI) (CA INDEX NAME)



RN 310442-94-5 CAPLUS

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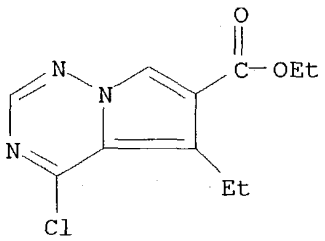
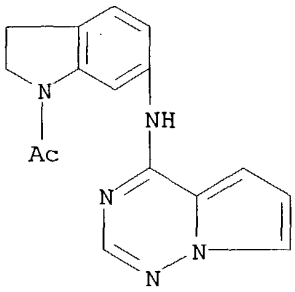
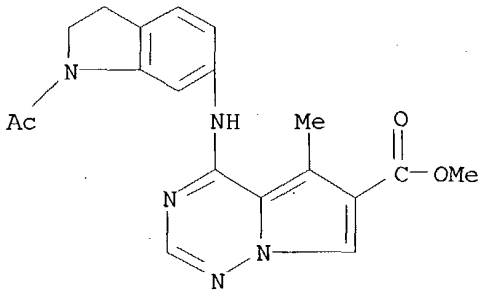


IT **310442-23-0P 310442-57-0P 310442-60-5P**
310442-72-9P 310442-75-2P 310442-77-4P
310442-79-6P
RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(preparation of pyrrolotriazines as kinases inhibitors useful in treating inflammation, cancer, and proliferative diseases)

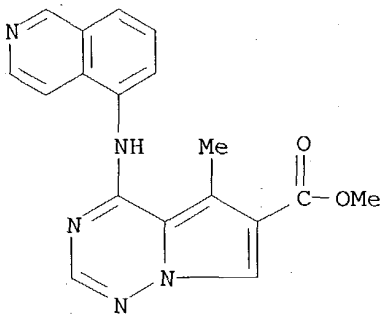
RN 310442-23-0 CAPLUS
CN 1H-Indol-6-amine, 1-acetyl-2,3-dihydro-N-pyrrolo[2,1-f][1,2,4]triazin-4-yl- (9CI) (CA INDEX NAME)



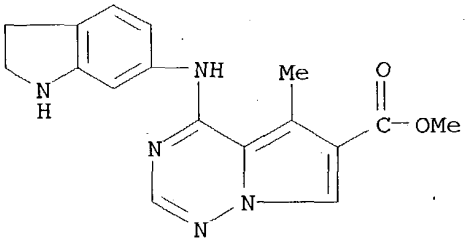
RN 310442-57-0 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 4-[(1-acetyl-2,3-dihydro-1H-indol-6-yl)amino]-5-methyl-, methyl ester (9CI) (CA INDEX NAME)



RN 310442-60-5 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 4-(5-isoquinolinylamino)-5-methyl-, methyl ester (9CI) (CA INDEX NAME)



RN 310442-72-9 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 4-[(2,3-dihydro-1H-indol-6-yl)amino]-5-methyl-, methyl ester (9CI) (CA INDEX NAME)



RN 310442-75-2 CAPLUS
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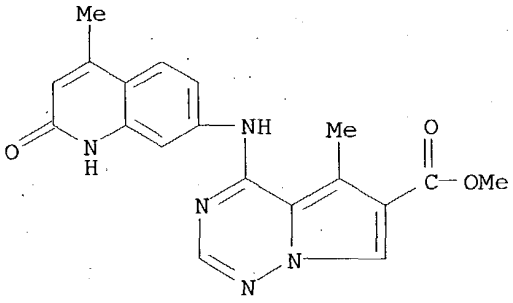
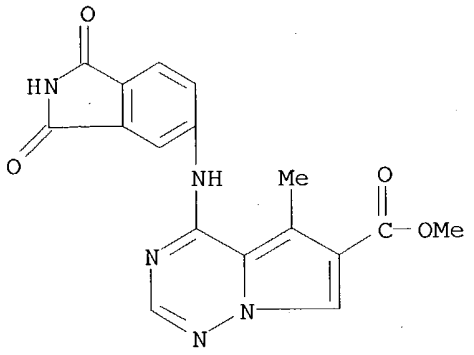


RN 310442-77-4 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 4-[(2,3-dihydro-1,3-dioxo-1H-isoindol-5-yl)amino]-5-methyl-, methyl ester (9CI) (CA INDEX NAME)



RN 310442-79-6 CAPLUS
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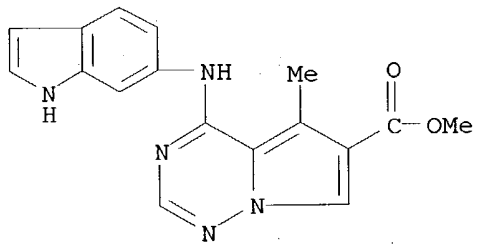
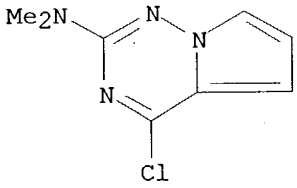


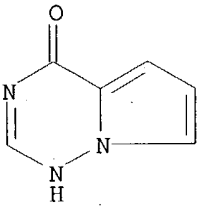
IT **175726-62-2**
RL: RCT (Reactant); RACT (Reactant or reagent)
(preparation of pyrrolotriazines as kinases inhibitors useful in treating inflammation, cancer, and proliferative diseases)
RN 175726-62-2 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazin-2-amine, 4-chloro-N,N-dimethyl- (9CI) (CA INDEX NAME)



IT **159326-71-3P**, Pyrrolo[2,1-f][1,2,4]triazin-4(1H)-one
310430-81-0P 310430-94-5P 310430-97-8P
310431-16-4P 310431-29-9P 310435-15-5P
310436-48-7P 310436-60-3P 310444-78-1P
310444-86-1P 310444-87-2P 310444-88-3P
310444-89-4P 310444-90-7P 310444-95-2P
310444-96-3P 310452-44-9P
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(preparation of pyrrolotriazines as kinases inhibitors useful in treating inflammation, cancer, and proliferative diseases)

RN 159326-71-3 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazin-4(1H)-one (9CI) (CA INDEX NAME)



RN 310430-81-0 CAPLUS
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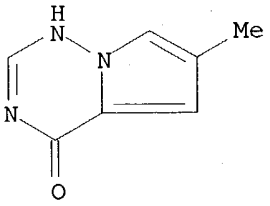

RN 310430-94-5 CAPLUS
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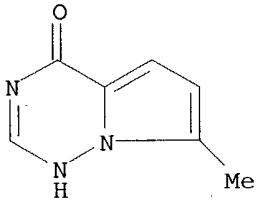

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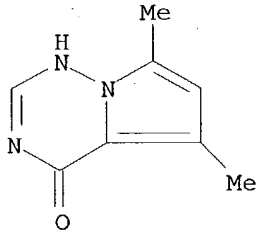

RN 310431-16-4 CAPLUS
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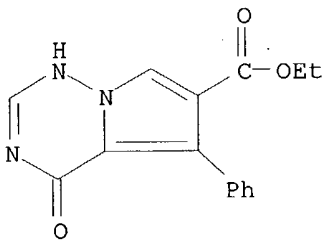
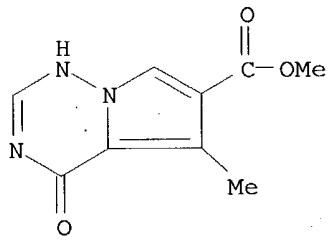
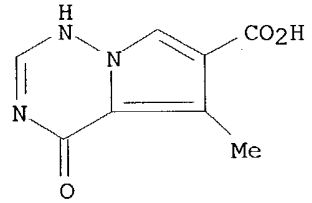
phenyl-, ethyl ester (9CI) (CA INDEX NAME)



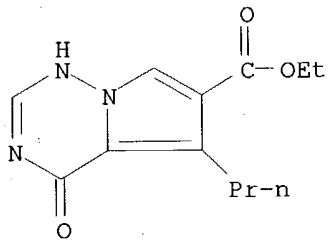
RN 310431-29-9 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-5-methyl-4-oxo-, methyl ester (9CI) (CA INDEX NAME)



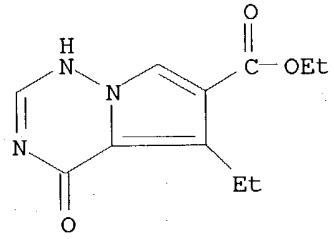
RN 310435-15-5 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-5-methyl-4-oxo- (9CI) (CA INDEX NAME)



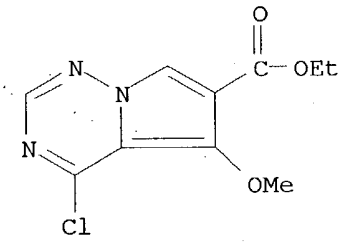
RN 310436-48-7 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-4-oxo-5-propyl-, ethyl ester (9CI) (CA INDEX NAME)



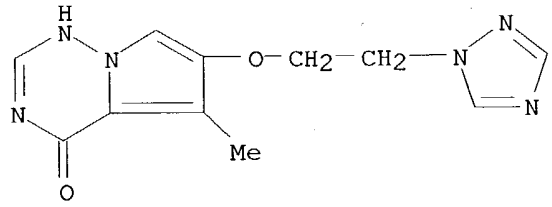
RN 310436-60-3 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 5-ethyl-1,4-dihydro-4-oxo-, ethyl ester (9CI) (CA INDEX NAME)



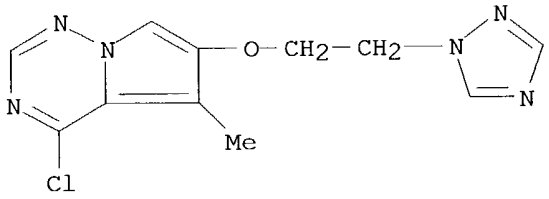
RN 310444-78-1 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 4-chloro-5-methoxy-, ethyl ester (9CI) (CA INDEX NAME)



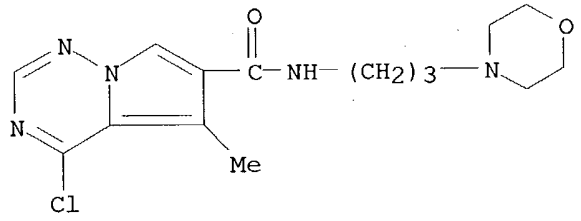
RN 310444-86-1 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazin-4(1H)-one, 5-methyl-6-[2-(1H-1,2,4-triazol-1-yl)ethoxy]- (9CI) (CA INDEX NAME)



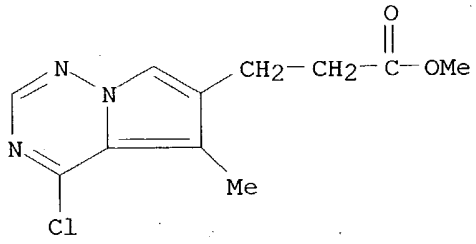
RN 310444-87-2 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine, 4-chloro-5-methyl-6-[2-(1H-1,2,4-triazol-1-yl)ethoxy]- (9CI) (CA INDEX NAME)



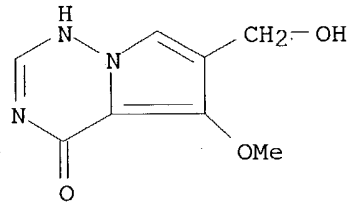
RN 310444-88-3 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxamide, 4-chloro-5-methyl-N-[3-(4-morpholinyl)propyl]- (9CI) (CA INDEX NAME)



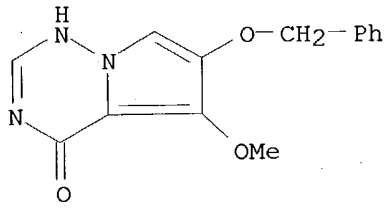
RN 310444-89-4 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-propanoic acid, 4-chloro-5-methyl-, methyl ester (9CI) (CA INDEX NAME)



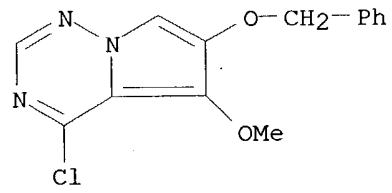
RN 310444-90-7 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazin-4(1H)-one, 6-(hydroxymethyl)-5-methoxy- (9CI) (CA INDEX NAME)



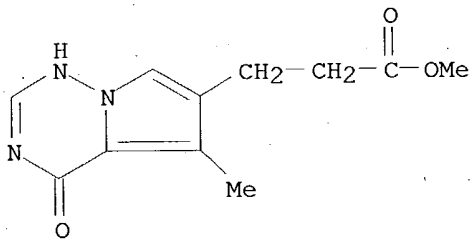
RN 310444-95-2 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazin-4(1H)-one, 5-methoxy-6-(phenylmethoxy)- (9CI) (CA INDEX NAME)



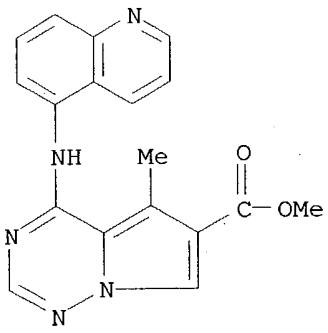
RN 310444-96-3 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine, 4-chloro-5-methoxy-6-(phenylmethoxy)- (9CI) (CA INDEX NAME)



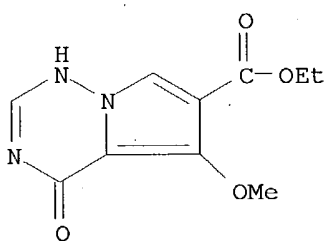
RN 310452-44-9 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-propanoic acid, 1,4-dihydro-5-methyl-4-oxo-, methyl ester (9CI) (CA INDEX NAME)



IT **310443-48-2P 310443-54-0P**
RL: SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(preparation of pyrrolotriazines as kinases inhibitors useful in treating inflammation, cancer, and proliferative diseases)
RN 310443-48-2 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 5-methyl-4-(5-quinolinylamino)-, methyl ester (9CI) (CA INDEX NAME)



RN 310443-54-0 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-5-methoxy-4-oxo-, ethyl ester (9CI) (CA INDEX NAME)



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L7 17 L3

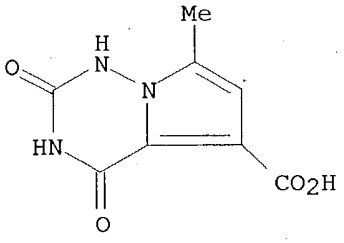
=> s l7 not l4 not l5
L8 15 L7 NOT L4 NOT L5

=> sort py l8
SORT ENTIRE ANSWER SET? (Y)/N:.
PROCESSING COMPLETED FOR L8
L9 15 SORT L8 PY

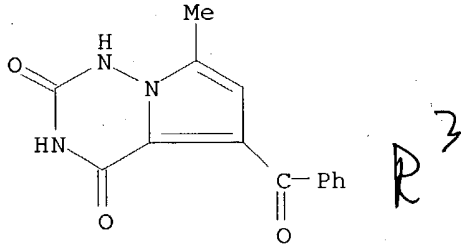
=> d 1-15 ibib pi hitstr

L9 ANSWER 1 OF 15 CAPLUS COPYRIGHT 2004 ACS on STN
ACCESSION NUMBER: 1979:611372 CAPLUS
DOCUMENT NUMBER: 91:211372
TITLE: Synthesis of a new bridgehead nitrogen heterocyclic system. Pyrrolo[2,1-f]-1,2,4-triazine derivatives
AUTHOR(S): Migliara, Onofrio; Petruso, Salvatore; Sprio, Vincenzo
CORPORATE SOURCE: Fac. Farmacia, Univ. Palermo, Palermo, 90123, Italy
SOURCE: Journal of Heterocyclic Chemistry (1979), 16(5), 833-4
CODEN: JHTCAD; ISSN: 0022-152X
DOCUMENT TYPE: Journal
LANGUAGE: English
OTHER SOURCE(S): CASREACT 91:211372

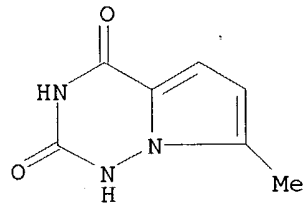
IT **71971-29-4P**
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(preparation and pyrolysis of)
RN 71971-29-4 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-5-carboxylic acid, 1,2,3,4-tetrahydro-7-methyl-2,4-dioxo- (9CI) (CA INDEX NAME)



IT **71971-30-7P 71971-31-8P**
RL: SPN (Synthetic preparation); PREP (Preparation)
(preparation of)
RN 71971-30-7 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-2,4(1H,3H)-dione, 5-benzoyl-7-methyl- (9CI) (CA INDEX NAME)

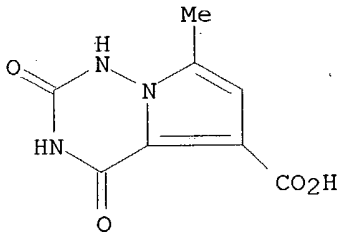


RN 71971-31-8 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-2,4(1H,3H)-dione, 7-methyl- (9CI) (CA INDEX NAME)

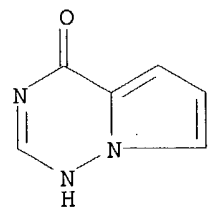


L9 ANSWER 2 OF 15 CAPLUS COPYRIGHT 2004 ACS on STN
ACCESSION NUMBER: 1983:143216 CAPLUS
DOCUMENT NUMBER: 98:143216
TITLE: Carbon-13 NMR characterization of carboxyl derivatives of 1-ureidopyrroles

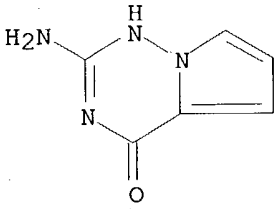
AUTHOR(S): Lamartina, Liliana; Migliara, Onofrio; Sprio, Vincenzo
CORPORATE SOURCE: Fac. Farm., Univ. Palermo, Palermo, 90123, Italy
SOURCE: Journal of Heterocyclic Chemistry (1982), 19(6), 1381-4
CODEN: JHTCAD; ISSN: 0022-152X
DOCUMENT TYPE: Journal
LANGUAGE: English
IT **71971-29-4P**
RL: SPN (Synthetic preparation); PREP (Preparation)
(preparation of)
RN 71971-29-4 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-5-carboxylic acid, 1,2,3,4-tetrahydro-7-methyl-2,4-dioxo- (9CI) (CA INDEX NAME)



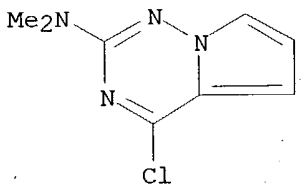
L9 ANSWER 3 OF 15 CAPLUS COPYRIGHT 2004 ACS on STN
ACCESSION NUMBER: 1995:51452 CAPLUS
DOCUMENT NUMBER: 122:9999
TITLE: Synthesis of pyrrolo[2,1-f][1,2,4]triazine congeners of nucleic acid purines via the N-amination of 2-substituted pyrroles
AUTHOR(S): Patil, Shirish A.; Otter, Brian A.; Klein, Robert S.
CORPORATE SOURCE: Albert Einstein Coll., Medicine Cancer Cent., Bronx, NY, 10467, USA
SOURCE: Journal of Heterocyclic Chemistry (1994), 31(4), 781-6
CODEN: JHTCAD; ISSN: 0022-152X
DOCUMENT TYPE: Journal
LANGUAGE: English
OTHER SOURCE(S): CASREACT 122:9999
IT **159326-71-3P**, Pyrrolo[2,1-f][1,2,4]triazin-4(1H)-one
159326-75-7P
RL: SPN (Synthetic preparation); PREP (Preparation)
(synthesis of pyrrolotriazine congeners of nucleic acid purines via amination of pyrroles)
RN 159326-71-3 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazin-4(1H)-one (9CI) (CA INDEX NAME)



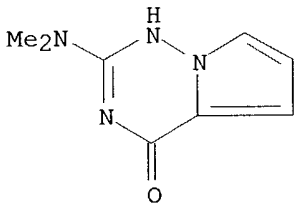
RN 159326-75-7 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazin-4(1H)-one, 2-amino- (9CI) (CA INDEX NAME)



L9 ANSWER 4 OF 15 CAPLUS COPYRIGHT 2004 ACS on STN
ACCESSION NUMBER: 1996:134791 CAPLUS
DOCUMENT NUMBER: 124:289464
TITLE: A ready one-pot preparation for pyrrolo[2,1-f][1,2,4]triazine and pyrazolo[5,1-c]pyrimido[4,5-e][1,2,4]triazine derivatives
AUTHOR(S): Quintela, Jose M.; Moreira, Maria J.; Peinador, Carlos
CORPORATE SOURCE: Facultad Ciencias, Univ. La Coruna, La Coruna, E-15071, Spain
SOURCE: Tetrahedron (1996), 52(8), 3037-48
CODEN: TETRAB; ISSN: 0040-4020
PUBLISHER: Elsevier
DOCUMENT TYPE: Journal
LANGUAGE: English
IT **175726-62-2P**
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(preparation of pyrrolo- and pyrazolopyrimidotriazines)
RN 175726-62-2 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazin-2-amine, 4-chloro-N,N-dimethyl- (9CI) (CA INDEX NAME)

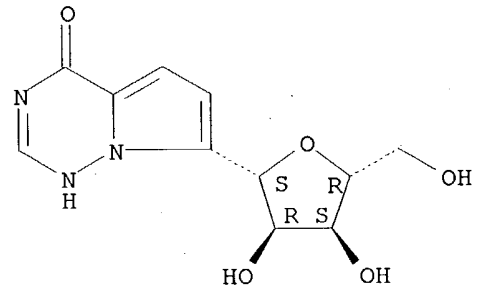


IT **175726-72-4P**
RL: SPN (Synthetic preparation); PREP (Preparation)
(preparation of pyrrolo- and pyrazolopyrimidotriazines)
RN 175726-72-4 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazin-4(1H)-one, 2-(dimethylamino)- (9CI) (CA INDEX NAME)



L9 ANSWER 5 OF 15 CAPLUS COPYRIGHT 2004 ACS on STN
ACCESSION NUMBER: 1996:109112 CAPLUS
DOCUMENT NUMBER: 124:290158
TITLE: Conformational properties of purine-like C-nucleosides
AUTHOR(S): Otter, Brian A.; Klein, Robert S.
CORPORATE SOURCE: Dep. of Oncology, Montefiore Medical Center, Bronx, NY, 10467, USA
SOURCE: Nucleosides & Nucleotides (1996), 15(1-3), 793-807
CODEN: NUNUD5; ISSN: 0732-8311
PUBLISHER: Dekker
DOCUMENT TYPE: Journal
LANGUAGE: English
IT **175688-18-3**
RL: PRP (Properties)
(conformation and hydrogen bond of purine-like C-nucleosides)
RN 175688-18-3 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazin-4(1H)-one, 7-β-D-ribofuranosyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



L9 ANSWER 6 OF 15 CAPLUS COPYRIGHT 2004 ACS on STN
ACCESSION NUMBER: 2001:152684 CAPLUS
DOCUMENT NUMBER: 134:193452
TITLE: Preparation of pyrrolotriazine derivatives as secretory phospholipase A2 (sPLA2) inhibitors
INVENTOR(S): Ohtani, Mitsuaki; Fuji, Masahiro; Ogawa, Tomoyuki
PATENT ASSIGNEE(S): Shionogi & Co., Ltd., Japan
SOURCE: PCT Int. Appl., 80 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: Japanese
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2001014378	A1	20010301	WO 2000-JP5357	20000810

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM

RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG

PRIORITY APPLN. INFO.: JP 1999-235957 A 19990823

OTHER SOURCE(S): MARPAT 134:193452

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI WO 2001014378	A1	20010301	WO 2000-JP5357	20000810

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM

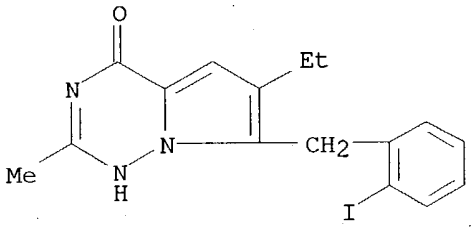
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG

IT **327976-40-9**

RL: RCT (Reactant); RACT (Reactant or reagent)
(preparation of pyrrolotriazine derivs. as secretory phospholipase A2 (sPLA2) inhibitors)

RN 327976-40-9 CAPLUS

CN Pyrrolo[2,1-f][1,2,4]triazin-4(1H)-one, 6-ethyl-7-[(2-iodophenyl)methyl]-2-methyl- (9CI) (CA INDEX NAME)

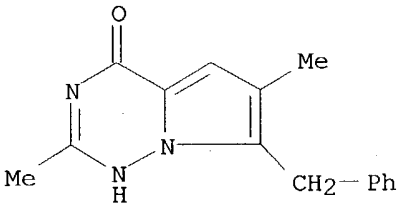


IT **327976-14-7P 327976-16-9P 327976-30-7P 327976-32-9P 327976-36-3P**

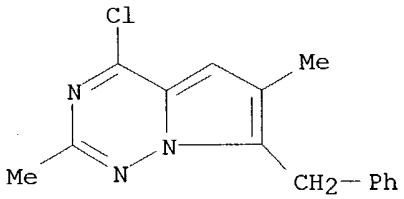
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(preparation of pyrrolotriazine derivs. as secretory phospholipase A2 (sPLA2) inhibitors)

RN 327976-14-7 CAPLUS

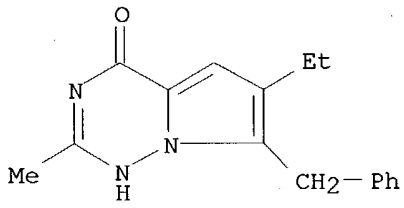
CN Pyrrolo[2,1-f][1,2,4]triazin-4(1H)-one, 2,6-dimethyl-7-(phenylmethyl)- (9CI) (CA INDEX NAME)



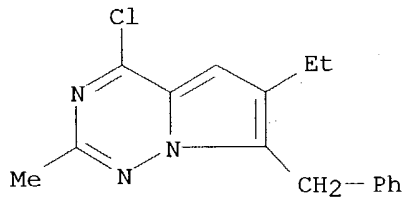
RN 327976-16-9 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine, 4-chloro-2,6-dimethyl-7-(phenylmethyl)- (9CI) (CA INDEX NAME)



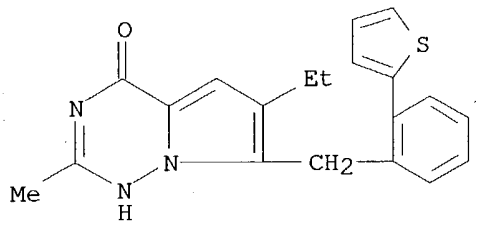
RN 327976-30-7 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazin-4(1H)-one, 6-ethyl-2-methyl-7-(phenylmethyl)- (9CI) (CA INDEX NAME)



RN 327976-32-9 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine, 4-chloro-6-ethyl-2-methyl-7-(phenylmethyl)- (9CI) (CA INDEX NAME)



RN 327976-36-3 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazin-4(1H)-one, 6-ethyl-2-methyl-7-[[2-(2-thienyl)phenyl]methyl]- (9CI) (CA INDEX NAME)



REFERENCE COUNT: 6 THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L9 ANSWER 7 OF 15 CAPLUS COPYRIGHT 2004 ACS on STN
ACCESSION NUMBER: 2002:391720 CAPLUS
DOCUMENT NUMBER: 136:386144
TITLE: Preparation of pyrrolo[2,1-f][1,2,4]triazine carboxylic acid derivatives for use in treating p38 kinase-associated conditions
INVENTOR(S): Leftheris, Katerina; Barrish, Joel; Hynes, John; Wrobleski, Stephen T.
PATENT ASSIGNEE(S): Bristol-Myers Squibb Company, USA
SOURCE: PCT Int. Appl., 108 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 2
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2002040486	A2	20020523	WO 2001-US49982	20011107
WO 2002040486	A3	20030912		
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
AU 2002032760	A5	20020527	AU 2002-32760	20011107
EE 200300227	A	20031015	EE 2003-227	20011107
EP 1363910	A2	20031126	EP 2001-992298	20011107
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR				
NO 2003002229	A	20030716	NO 2003-2229	20030516

PRIORITY APPLN. INFO.:
US 2000-249877P P 20001117
US 2001-310561P P 20010807
WO 2001-US49982 W 20011107

OTHER SOURCE(S): MARPAT 136:386144

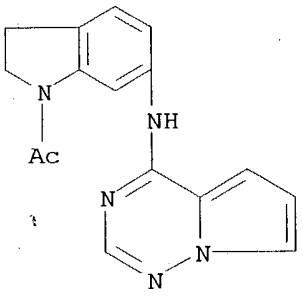
	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2002040486	A2	20020523	WO 2001-US49982	20011107
	WO 2002040486	A3	20030912		
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN,				

CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG

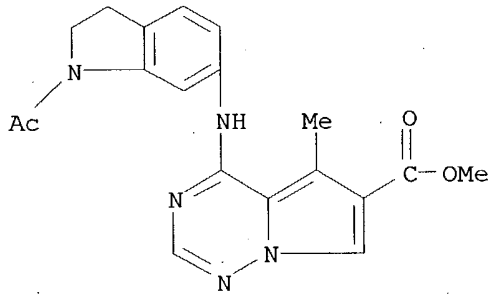
AU 2002032760	A5	20020527	AU 2002-32760	20011107
EE 200300227	A	20031015	EE 2003-227	20011107
EP 1363910	A2	20031126	EP 2001-992298	20011107
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR				
NO 2003002229	A	20030716	NO 2003-2229	20030516

IT **310442-23-0P**, 1-[2,3-Dihydro-6-[pyrrolo[2,1-f][1,2,4]triazin-4-ylamino]-1H-indol-1-yl]ethanone **310442-57-0P**, 4-[[1-Acetyl-2,3-dihydro-1H-indol-6-yl]amino]-5-methylpyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid methyl ester
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(drug; preparation of pyrrolo[2,1-f][1,2,4]triazine carboxylic acid derivs. for use in treating p38 kinase-associated conditions)

RN 310442-23-0 CAPLUS
CN 1H-Indol-6-amine, 1-acetyl-2,3-dihydro-N-pyrrolo[2,1-f][1,2,4]triazin-4-yl- (9CI) (CA INDEX NAME)



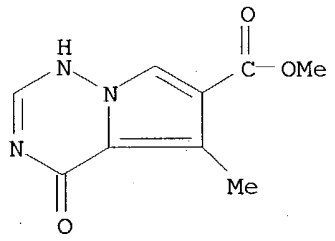
RN 310442-57-0 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 4-[(1-acetyl-2,3-dihydro-1H-indol-6-yl)amino]-5-methyl-, methyl ester (9CI) (CA INDEX NAME)



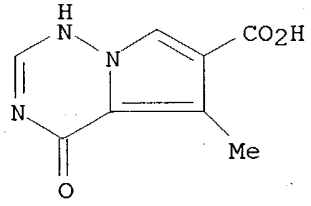
IT **310431-29-9P 310435-15-5P 310442-40-1P**,

4-Chloro-5-methylpyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid methyl ester **310443-54-0P**, 4-Hydroxy-5-methoxypyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid ethyl ester **310444-88-3P**, 4-Chloro-5-methyl-N-[3-[4-morpholinyl]propyl]pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid amide **310444-89-4P**, 4-Chloro-5-methylpyrrolo[2,1-f][1,2,4]triazine-6-propanoic acid methyl ester **310444-90-7P 310444-95-2P 310444-96-3P**, 4-Chloro-5-methoxy-6-[phenylmethoxy]pyrrolo[2,1-f][1,2,4]triazine **310452-44-9P**, 4-Hydroxy-5-methylpyrrolo[2,1-f][1,2,4]triazine-6-propanoic acid methyl ester **427878-41-9P 427878-70-4P**
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(intermediate; preparation of pyrrolo[2,1-f][1,2,4]triazine carboxylic acid derivs. for use in treating p38 kinase-associated conditions)

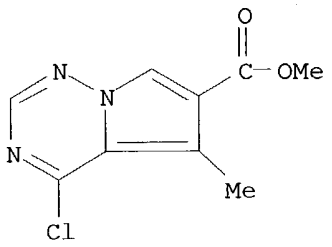
RN 310431-29-9 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-5-methyl-4-oxo-, methyl ester (9CI) (CA INDEX NAME)



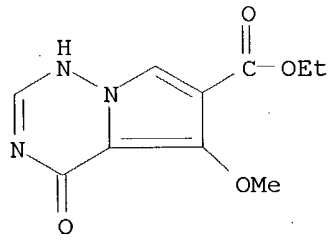
RN 310435-15-5 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-5-methyl-4-oxo- (9CI) (CA INDEX NAME)



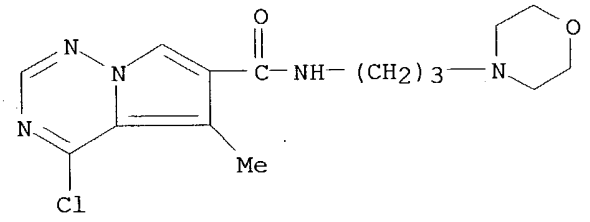
RN 310442-40-1 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 4-chloro-5-methyl-, methyl ester (9CI) (CA INDEX NAME)



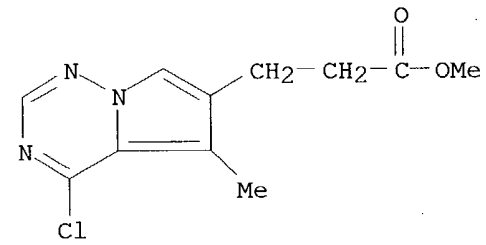
RN 310443-54-0 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-5-methoxy-4-oxo-, ethyl ester (9CI) (CA INDEX NAME)



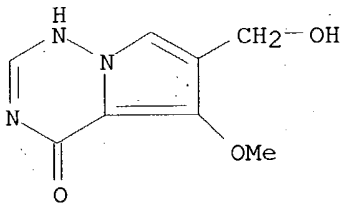
RN 310444-88-3 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxamide, 4-chloro-5-methyl-N-[3-(4-morpholinyl)propyl]- (9CI) (CA INDEX NAME)



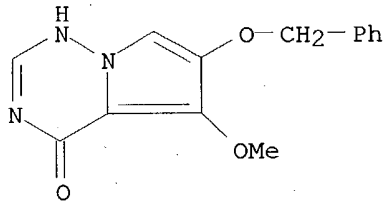
RN 310444-89-4 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-propanoic acid, 4-chloro-5-methyl-, methyl ester (9CI) (CA INDEX NAME)



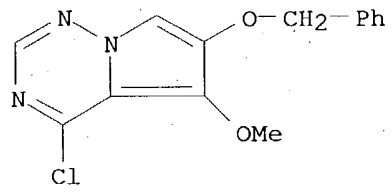
RN 310444-90-7 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazin-4(1H)-one, 6-(hydroxymethyl)-5-methoxy- (9CI) (CA INDEX NAME)



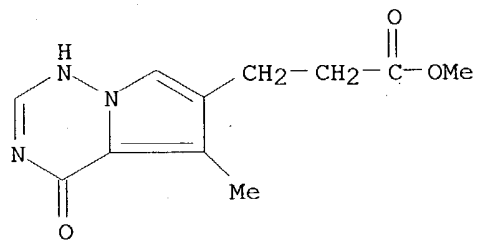
RN 310444-95-2 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazin-4(1H)-one, 5-methoxy-6-(phenylmethoxy)- (9CI) (CA INDEX NAME)



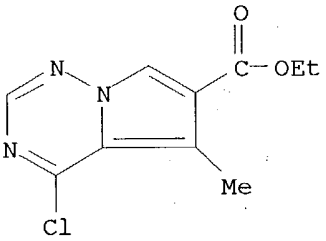
RN 310444-96-3 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine, 4-chloro-5-methoxy-6-(phenylmethoxy)- (9CI) (CA INDEX NAME)



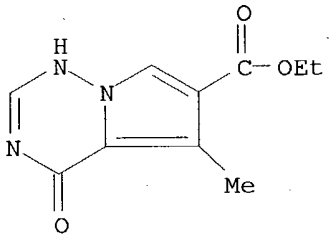
RN 310452-44-9 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-propanoic acid, 1,4-dihydro-5-methyl-4-oxo-, methyl ester (9CI) (CA INDEX NAME)



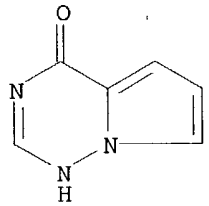
RN 427878-41-9 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 4-chloro-5-methyl-, ethyl ester (9CI) (CA INDEX NAME)



RN 427878-70-4 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-5-methyl-4-oxo-, ethyl ester (9CI) (CA INDEX NAME)



IT **159326-71-3**, Pyrrolo[2,1-f][1,2,4]triazin-4(1H)-one
RL: RCT (Reactant); RACT (Reactant or reagent)
(reactant; preparation of pyrrolo[2,1-f][1,2,4]triazine carboxylic acid derivs. for use in treating p38 kinase-associated conditions)
RN 159326-71-3 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazin-4(1H)-one (9CI) (CA INDEX NAME)



L9 ANSWER 8 OF 15 CAPLUS COPYRIGHT 2004 ACS on STN
ACCESSION NUMBER: 2003:950844 CAPLUS
DOCUMENT NUMBER: 140:5075
TITLE: Pyrrolotriazinone compounds and their use to treat diseases
INVENTOR(S): Lombardo, Louis J.; Bhide, Rajeev S.; Kim, Kyoung S.; Lu, Songfeng

PATENT ASSIGNEE(S): Bristol-Myers Squibb Company, USA
SOURCE: PCT Int. Appl., 106 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2003099286	A1	20031204	WO 2003-US16179	20030520

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM

RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG

US 2003232832	A1	20031218	US 2003-441848	20030520

PRIORITY APPLN. INFO.: US 2002-382197P P 20020521
OTHER SOURCE(S): MARPAT 140:5075

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2003099286	A1	20031204	WO 2003-US16179	20030520

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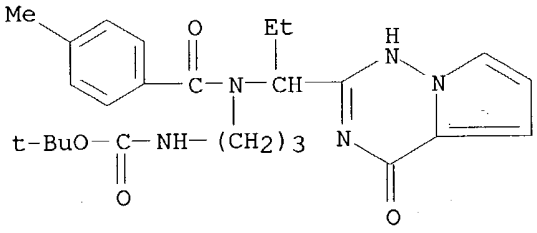
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG

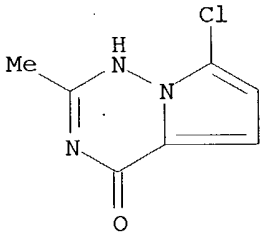
US 2003232832	A1	20031218	US 2003-441848	20030520

IT **628733-89-1P 628734-14-5P 628734-24-7P**
628734-34-9P 628734-46-3P
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(intermediates; in preparation of pyrrolotriazinone compds. useful for inducing mitotic arrest, anticancer agents, and other disease treatment)

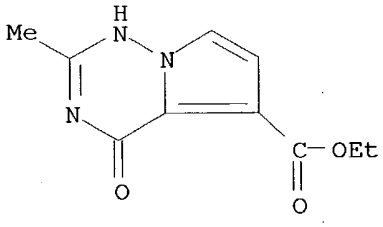
RN 628733-89-1 CAPLUS
CN Carbamic acid, [3-[[1-(1,4-dihydro-4-oxopyrrolo[2,1-f][1,2,4]triazin-2-yl)propyl](4-methylbenzoyl)amino]propyl]-, 1,1-dimethylethyl ester (9CI) (CA INDEX NAME)



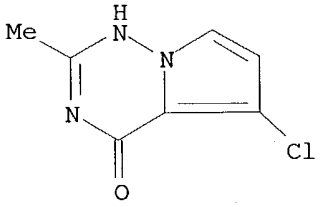
RN 628734-14-5 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazin-4(1H)-one, 7-chloro-2-methyl- (9CI) (CA INDEX NAME)



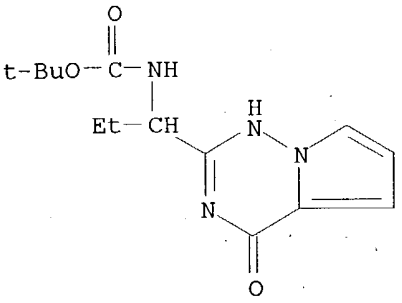
RN 628734-24-7 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-5-carboxylic acid, 1,4-dihydro-2-methyl-4-oxo-, ethyl ester (9CI) (CA INDEX NAME)



RN 628734-34-9 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazin-4(1H)-one, 5-chloro-2-methyl- (9CI) (CA INDEX NAME)



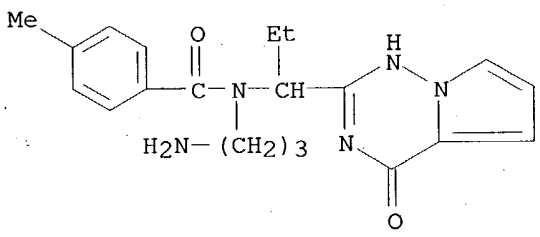
RN 628734-46-3 CAPLUS
CN Carbamic acid, [1-(1,4-dihydro-4-oxopyrrolo[2,1-f][1,2,4]triazin-2-yl)propyl]-, 1,1-dimethylethyl ester (9CI) (CA INDEX NAME)



IT **628733-07-3P 628733-41-5P**
RL: SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(preparation of pyrrolotriazinone compds. useful for inducing mitotic arrest, anticancer agents, and other disease treatment)
RN 628733-07-3 CAPLUS
CN Benzamide, N-(3-aminopropyl)-N-[1-(1,4-dihydro-4-oxopyrrolo[2,1-f][1,2,4]triazin-2-yl)propyl]-4-methyl-, mono(trifluoroacetate) (9CI) (CA INDEX NAME)

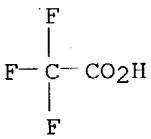
CM 1

CRN 628733-06-2
CMF C20 H25 N5 O2

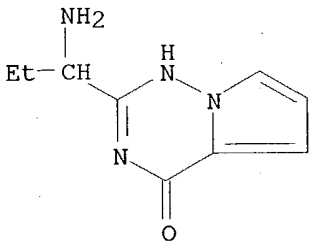


CM 2

CRN 76-05-1
CMF C2 H F3 O2



RN 628733-41-5 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazin-4(1H)-one, 2-(1-aminopropyl)-, monohydrochloride (9CI) (CA INDEX NAME)



● HCl

REFERENCE COUNT: 1 THERE ARE 1 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L9 ANSWER 9 OF 15 CAPLUS COPYRIGHT 2004 ACS on STN
ACCESSION NUMBER: 2003:875265 CAPLUS
DOCUMENT NUMBER: 139:364963
TITLE: Aryl ketone pyrrolo-triazine compounds useful as kinase inhibitors, particularly p38 kinases, and their preparation, pharmaceutical compositions, and use
INVENTOR(S): Dyckman, Alaric; Leftheris, Katerina; Hynes, John
PATENT ASSIGNEE(S): Bristol-Myers Squibb Company, USA
SOURCE: PCT Int. Appl., 45 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2003091229	A1	20031106	WO 2003-US12420	20030418
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
US 2003232831	A1	20031218	US 2003-420445	20030422

PRIORITY APPLN. INFO.: US 2002-374907P P 20020423
OTHER SOURCE(S): MARPAT 139:364963

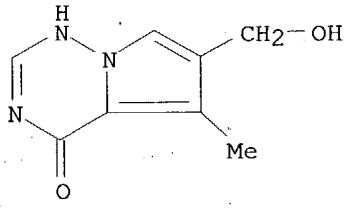
	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2003091229	A1	20031106	WO 2003-US12420	20030418
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR,				

LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG

US 2003232831 A1 20031218 US 2003-420445 20030422

IT **621685-54-9P 621685-55-0P 621685-56-1P**
621685-57-2P 621685-58-3P 621685-59-4P
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(intermediate; preparation of aryl ketone pyrrolotriazine compds. as p38 kinase inhibitors)

RN 621685-54-9 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazin-4(1H)-one, 6-(hydroxymethyl)-5-methyl- (9CI) (CA INDEX NAME)



RN 621685-55-0 CAPLUS
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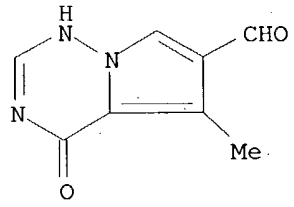
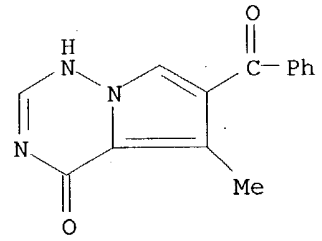
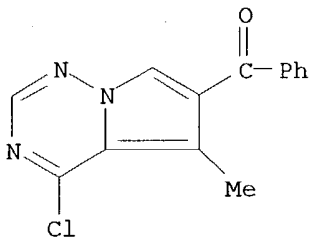


RN 621685-56-1 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazin-4(1H)-one, 6-benzoyl-5-methyl- (9CI) (CA INDEX NAME)



RN 621685-57-2 CAPLUS
CN Methanone, (4-chloro-5-methylpyrrolo[2,1-f][1,2,4]triazin-6-yl)phenyl- (9CI) (CA INDEX NAME)



RN 621685-58-3 CAPLUS
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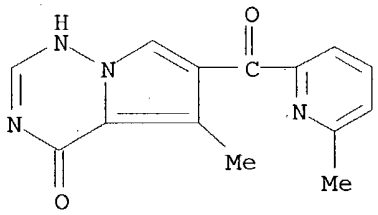
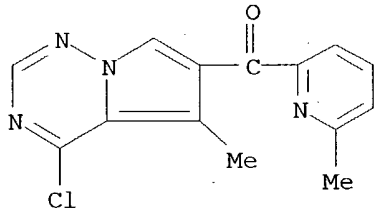
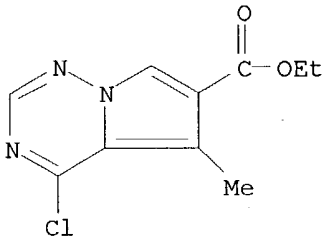


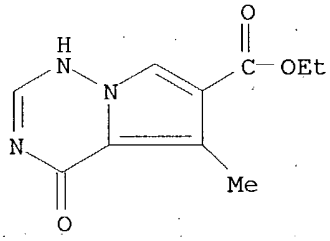
RN 621685-59-4 CAPLUS
CN Methanone, (4-chloro-5-methylpyrrolo[2,1-f][1,2,4]triazin-6-yl)(6-methyl-2-pyridinyl)- (9CI) (CA INDEX NAME)



IT **427878-41-9 427878-70-4**
RL: RCT (Reactant); RACT (Reactant or reagent)
(starting material; preparation of aryl ketone pyrrolotriazine compds. as p38 kinase inhibitors)
RN 427878-41-9 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 4-chloro-5-methyl-, ethyl ester (9CI) (CA INDEX NAME)



RN 427878-70-4 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-5-methyl-4-oxo-, ethyl ester (9CI) (CA INDEX NAME)



REFERENCE COUNT: 1 THERE ARE 1 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L9 ANSWER 10 OF 15 CAPLUS COPYRIGHT 2004 ACS on STN
ACCESSION NUMBER: 2003:875173 CAPLUS
DOCUMENT NUMBER: 139:381511
TITLE: Pyrrolotriazine aniline compounds useful as kinase inhibitors, particularly p38 kinases, and their preparation, pharmaceutical compositions, and use as antiinflammatory agents
INVENTOR(S): Dyckman, Alaric; Hynes, John; Leftheris, Katherina; Liu, Chunjian; Wrobleski, Stephen T.
PATENT ASSIGNEE(S): Bristol-Myers Squibb Company, USA
SOURCE: PCT Int. Appl., 158 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2003090912	A1	20031106	WO 2003-US12426	20030415
WO 2003090912	C2	20040108		

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ,

MD, RU, TJ, TM
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG

US 2004082582 A1 20040429 US 2003-420399 20030422
PRIORITY APPLN. INFO.: US 2002-374938P P 20020423
OTHER SOURCE(S): MARPAT 139:381511

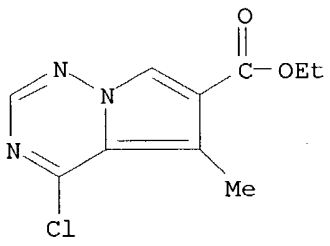
	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2003090912	A1	20031106	WO 2003-US12426	20030415
	WO 2003090912	C2	20040108		

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG

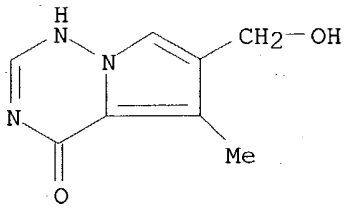
US 2004082582 A1 20040429 US 2003-420399 20030422

IT **427878-41-9P 621685-54-9P 621685-55-0P 621685-56-1P 621685-57-2P 621685-58-3P 621685-59-4P 623155-22-6P 623155-48-6P**
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(intermediate; preparation of pyrrolotriazine aniline compds. as p38 kinase inhibitors)

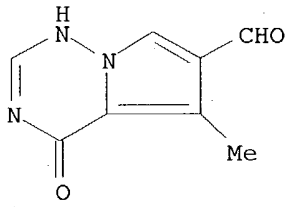
RN 427878-41-9 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 4-chloro-5-methyl-, ethyl ester (9CI) (CA INDEX NAME)



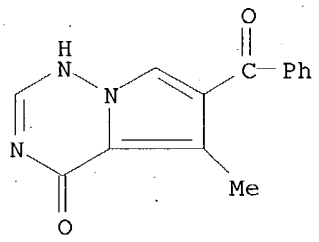
RN 621685-54-9 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazin-4(1H)-one, 6-(hydroxymethyl)-5-methyl- (9CI) (CA INDEX NAME)



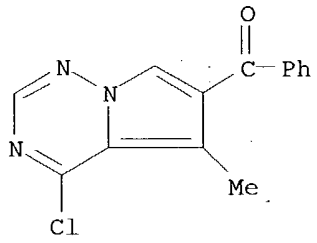
RN 621685-55-0 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxaldehyde, 1,4-dihydro-5-methyl-4-oxo- (9CI) (CA INDEX NAME)



RN 621685-56-1 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazin-4(1H)-one, 6-benzoyl-5-methyl- (9CI) (CA INDEX NAME)

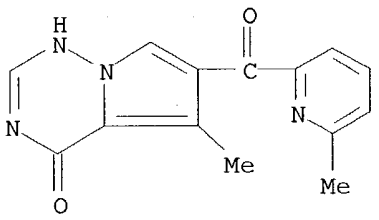


RN 621685-57-2 CAPLUS
CN Methanone, (4-chloro-5-methylpyrrolo[2,1-f][1,2,4]triazin-6-yl)phenyl- (9CI) (CA INDEX NAME)

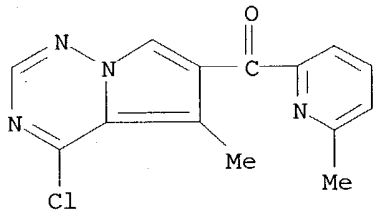


RN 621685-58-3 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazin-4(1H)-one, 5-methyl-6-[(6-methyl-2-

pyridinyl)carbonyl]- (9CI) (CA INDEX NAME)



RN 621685-59-4 CAPLUS
CN Methanone, (4-chloro-5-methylpyrrolo[2,1-f][1,2,4]triazin-6-yl)(6-methyl-2-pyridinyl)- (9CI) (CA INDEX NAME)



RN 623155-22-6 CAPLUS
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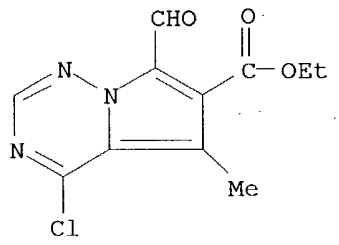

RN 623155-48-6 CAPLUS
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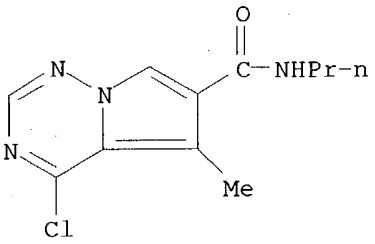
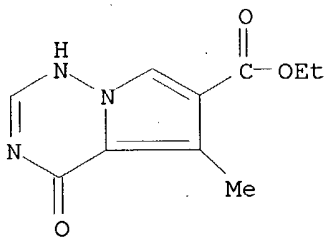


IT **427878-70-4**
RL: RCT (Reactant); RACT (Reactant or reagent)
(starting material; preparation of pyrrolotriazine aniline compds. as p38 kinase inhibitors)
RN 427878-70-4 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-5-methyl-4-oxo-, ethyl ester (9CI) (CA INDEX NAME)



REFERENCE COUNT: 5 THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L9 ANSWER 11 OF 15 CAPLUS COPYRIGHT 2004 ACS on STN
ACCESSION NUMBER: 2003:777390 CAPLUS
DOCUMENT NUMBER: 139:292275
TITLE: Methods for the preparation of pyrrolotriazine compounds useful as kinase inhibitors
INVENTOR(S): Godfrey, Jollie Duaine; Hynes, John; Dyckman, Alaric J.; Leftheris, Katerina; Shi, Zhongping; Wrobleski, Stephen T.; Doubleday, Wendel William; Grosso, John A.
PATENT ASSIGNEE(S): USA
SOURCE: U.S. Pat. Appl. Publ., 36 pp., Cont.-in-part of U.S. Ser. No. 36,293.
CODEN: USXXCO
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 2
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 2003186982	A1	20031002	US 2002-289010	20021106
US 2003069244	A1	20030410	US 2001-36293	20011107
US 6670357	B2	20031230		

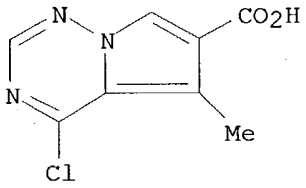
PRIORITY APPLN. INFO.: US 2000-249877P P 20001117

US 2001-310561P P 20010807
US 2001-36293 A2 20011107

OTHER SOURCE(S): MARPAT 139:292275

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	US 2003186982	A1	20031002	US 2002-289010	20021106
	US 2003069244	A1	20030410	US 2001-36293	20011107
	US 6670357	B2	20031230		

IT **607738-99-8**
RL: RCT (Reactant); RACT (Reactant or reagent)
(preparation of pyrrolotriazine derivative as kinase inhibitor)
RN 607738-99-8 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 4-chloro-5-methyl- (9CI) (CA INDEX NAME)



IT **427878-41-9P 427878-70-4P**
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(preparation of pyrrolotriazine derivative as kinase inhibitor)
RN 427878-41-9 CAPLUS
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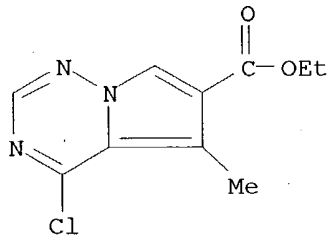
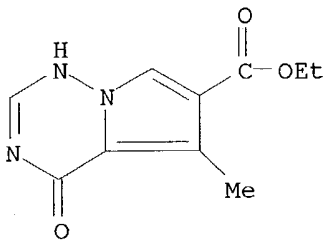


RN 427878-70-4 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-5-methyl-4-oxo-, ethyl ester (9CI) (CA INDEX NAME)



L9 ANSWER 12 OF 15 CAPLUS COPYRIGHT 2004 ACS on STN
ACCESSION NUMBER: 2003:396849 CAPLUS
DOCUMENT NUMBER: 138:401758
TITLE: Preparation of 5-substituted N-(1H-indazol-5-yl)pyrrolo[2,1-f][1,2,4]triazin-4-amines as antiproliferative agents
INVENTOR(S): Mastalerz, Harold; Zhang, Guifen; Tarrant, James G.; Vite, Gregory D.
PATENT ASSIGNEE(S): Bristol-Myers Squibb Company, USA
SOURCE: PCT Int. Appl., 74 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2003042172	A2	20030522	WO 2002-US36528	20021112
WO 2003042172	A3	20040129		

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM

RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG

US 2003186983	A1	20031002	US 2002-294281	20021114

PRIORITY APPLN. INFO.: US 2001-333014P P 20011114
OTHER SOURCE(S): MARPAT 138:401758

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI WO 2003042172	A2	20030522	WO 2002-US36528	20021112
WO 2003042172	A3	20040129		

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM

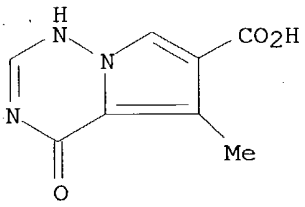
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG

US 2003186983 A1 20031002 US 2002-294281 20021114

IT **310435-15-5P**, 5-Methyl-4-oxo-3,4-dihydropyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid **529508-54-1P**, 5-Methyl-3H-pyrrolo[2,1-f][1,2,4]triazin-4-one **529508-56-3P**, 4-Chloro-5-methylpyrrolo[2,1-f][1,2,4]triazine **529508-57-4P**, 5-Bromomethyl-4-chloropyrrolo[2,1-f][1,2,4]triazine **529509-39-5P**, Acetic acid [[4-chloropyrrolo[2,1-f][1,2,4]triazin-5-yl]methyl] ester **529510-07-4P**, 4-Chloro-5-(2-methoxyethoxymethyl)pyrrolo[2,1-f][1,2,4]triazine
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(intermediate; preparation of N-(indazolyl)pyrrolotriazinamines as tyrosine kinase inhibitors for treatment of proliferative disorders and other diseases associated with signal transduction pathways)

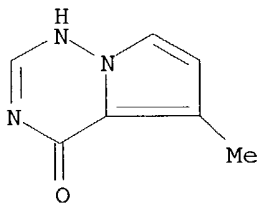
RN 310435-15-5 CAPLUS

CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-5-methyl-4-oxo- (9CI) (CA INDEX NAME)



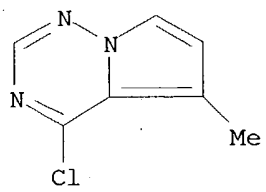
RN 529508-54-1 CAPLUS

CN Pyrrolo[2,1-f][1,2,4]triazin-4(1H)-one, 5-methyl- (9CI) (CA INDEX NAME)

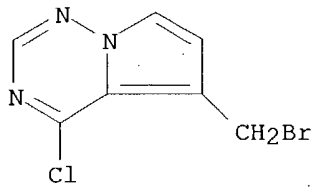


RN 529508-56-3 CAPLUS

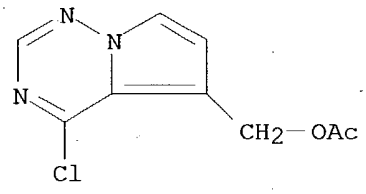
CN Pyrrolo[2,1-f][1,2,4]triazine, 4-chloro-5-methyl- (9CI) (CA INDEX NAME)



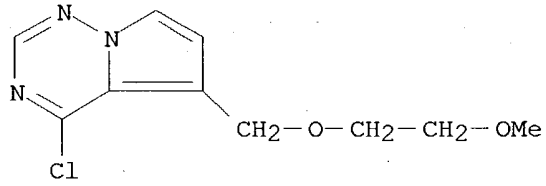
RN 529508-57-4 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine, 5-(bromomethyl)-4-chloro- (9CI) (CA INDEX NAME)



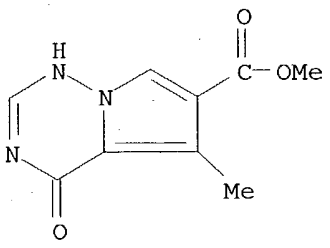
RN 529509-39-5 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-5-methanol, 4-chloro-, acetate (ester) (9CI) (CA INDEX NAME)



RN 529510-07-4 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine, 4-chloro-5-[(2-methoxyethoxy)methyl]- (9CI) (CA INDEX NAME)



IT **310431-29-9**, 5-Methyl-4-oxo-3,4-dihydropyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid methyl ester
RL: RCT (Reactant); RACT (Reactant or reagent)
(preparation of N-(indazolyl)pyrrolotriazinamines as tyrosine kinase inhibitors for treatment of proliferative disorders and other diseases associated with signal transduction pathways)
RN 310431-29-9 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-5-methyl-4-oxo-, methyl ester (9CI) (CA INDEX NAME)



L9 ANSWER 13 OF 15 CAPLUS COPYRIGHT 2004 ACS on STN
ACCESSION NUMBER: 2004:120859 CAPLUS
DOCUMENT NUMBER: 140:181471
TITLE: Preparation of pyrrolotriazines as tyrosine kinase activity inhibitors of growth factor receptors for the treatment of cancer
INVENTOR(S): Bhide, Rajeev S.; Borzilleri, Robert M.
PATENT ASSIGNEE(S): Bristol-Myers Squibb Company, USA
SOURCE: PCT Int. Appl., 71 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
	WO 2004013145	A1	20040212	WO 2003-US24273	20030804
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU				
	RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
	US 2004063708	A1	20040401	US 2003-633997	20030804

PRIORITY APPLN. INFO.: US 2002-400572P P 20020802
OTHER SOURCE(S): MARPAT 140:181471

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2004013145	A1	20040212	WO 2003-US24273	20030804
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU				
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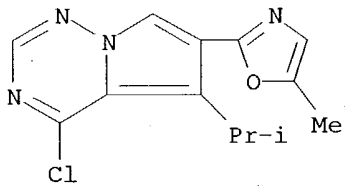
GW, ML, MR, NE, SN, TD, TG
US 2004063708 A1 20040401 US 2003-633997 20030804

IT **658084-81-2P**, 4-Chloro-5-(1-methylethyl)-6-(5-methyl-2-oxazolyl)pyrrolo[2,1-f][1,2,4]triazin
RL: PAC (Pharmacological activity); RCT (Reactant); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); RACT (Reactant or reagent); USES (Uses)
(drug candidate; preparation of pyrrolotriazines as tyrosine kinase activity inhibitors of growth factor receptors)

RN 658084-81-2 CAPLUS

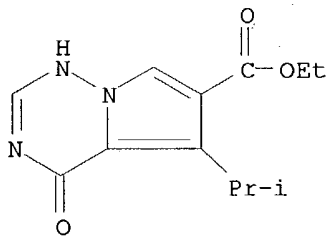
CN Pyrrolo[2,1-f][1,2,4]triazine, 4-chloro-5-(1-methylethyl)-6-(5-methyl-2-oxazolyl)- (9CI) (CA INDEX NAME)



IT **651744-40-0P**
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(drug candidate; preparation of pyrrolotriazines as tyrosine kinase activity inhibitors of growth factor receptors)

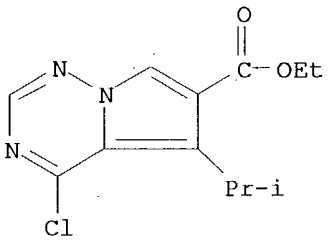
RN 651744-40-0 CAPLUS

CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-5-(1-methylethyl)-4-oxo-, ethyl ester (9CI) (CA INDEX NAME)

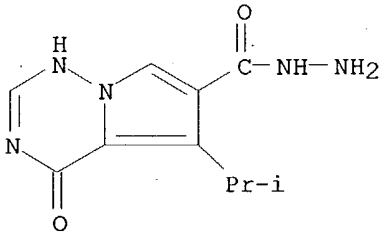


IT **658084-80-1P 658085-53-1P 658085-59-7P**
658085-60-0P 658085-61-1P 658085-62-2P
658085-63-3P 658085-64-4P 658085-65-5P,
6-Cyano-5-(1-methylethyl)pyrrolo[2,1-f][1,2,4]triazin-4(3H)-one
658085-66-6P 658085-67-7P, 5-(1-Methylethyl)-6-(1-methyl-1H-1,2,4-triazol-3-yl)pyrrolo[2,1-f][1,2,4]triazin-4(3H)-one
658085-69-9P 658085-70-2P 658085-71-3P,
4-Hydroxy-5-(1-methylethyl)pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid(2-oxopropyl)amide **658085-72-4P**
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(intermediate; preparation of pyrrolotriazines as tyrosine kinase activity inhibitors of growth factor receptors)

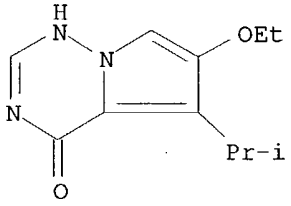
RN 658084-80-1 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 4-chloro-5-(1-methylethyl)-, ethyl ester (9CI) (CA INDEX NAME)



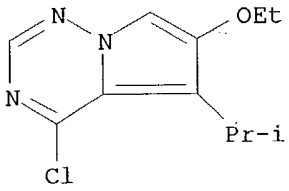
RN 658085-53-1 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-5-(1-methylethyl)-4-oxo-, hydrazide (9CI) (CA INDEX NAME)



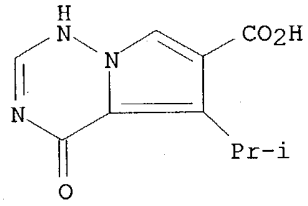
RN 658085-59-7 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazin-4(1H)-one, 6-ethoxy-5-(1-methylethyl)- (9CI) (CA INDEX NAME)



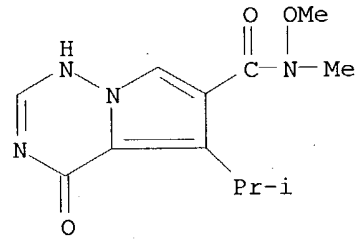
RN 658085-60-0 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine, 4-chloro-6-ethoxy-5-(1-methylethyl)- (9CI) (CA INDEX NAME)



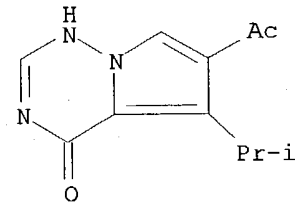
RN 658085-61-1 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-5-(1-methylethyl)-4-oxo- (9CI) (CA INDEX NAME)



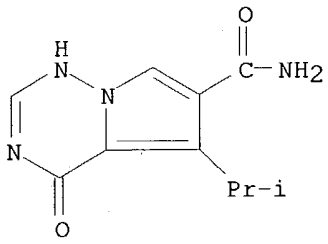
RN 658085-62-2 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxamide, 1,4-dihydro-N-methoxy-N-methyl-5-(1-methylethyl)-4-oxo- (9CI) (CA INDEX NAME)



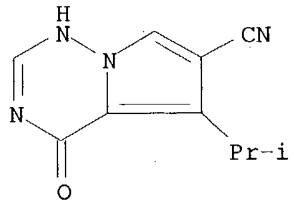
RN 658085-63-3 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazin-4(1H)-one, 6-acetyl-5-(1-methylethyl)- (9CI) (CA INDEX NAME)



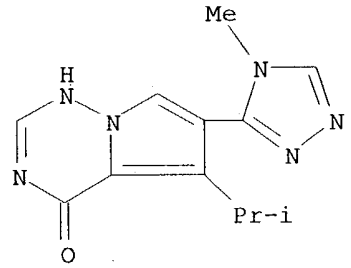
RN 658085-64-4 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxamide, 1,4-dihydro-5-(1-methylethyl)-4-oxo- (9CI) (CA INDEX NAME)



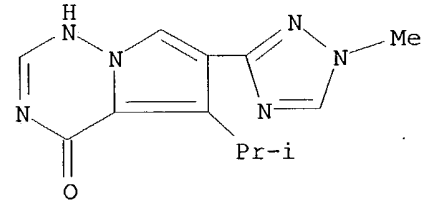
RN 658085-65-5 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carbonitrile, 1,4-dihydro-5-(1-methylethyl)-4-oxo- (9CI) (CA INDEX NAME)



RN 658085-66-6 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazin-4(1H)-one, 5-(1-methylethyl)-6-(4-methyl-4H-1,2,4-triazol-3-yl)- (9CI) (CA INDEX NAME)

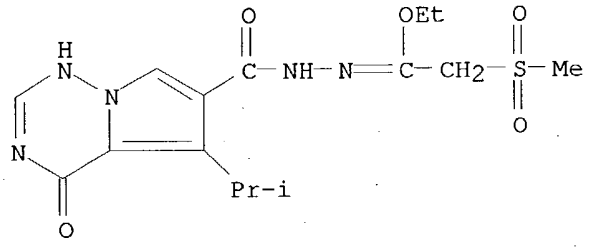


RN 658085-67-7 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazin-4(1H)-one, 5-(1-methylethyl)-6-(1-methyl-1H-1,2,4-triazol-3-yl)- (9CI) (CA INDEX NAME)

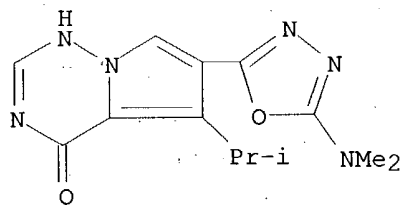


RN 658085-69-9 CAPLUS

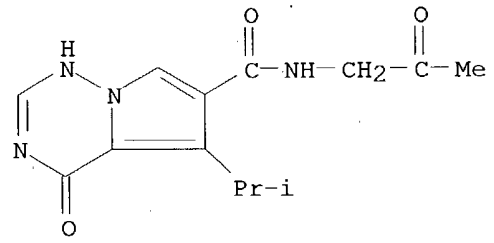
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-5-(1-methylethyl)-4-oxo-, [1-ethoxy-2-(methylsulfonyl)ethylidene]hydrazide (9CI) (CA INDEX NAME)



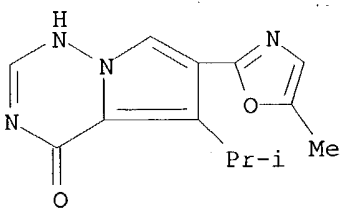
RN 658085-70-2 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazin-4(1H)-one, 6-[5-(dimethylamino)-1,3,4-oxadiazol-2-yl]-5-(1-methylethyl)- (9CI) (CA INDEX NAME)



RN 658085-71-3 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxamide, 1,4-dihydro-5-(1-methylethyl)-4-oxo-N-(2-oxopropyl)- (9CI) (CA INDEX NAME)



RN 658085-72-4 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazin-4(1H)-one, 5-(1-methylethyl)-6-(5-methyl-2-oxazolyl)- (9CI) (CA INDEX NAME)



L9 ANSWER 14 OF 15 CAPLUS COPYRIGHT 2004 ACS on STN
ACCESSION NUMBER: 2004:80698 CAPLUS
DOCUMENT NUMBER: 140:146173
TITLE: Preparation of pyrrolotriazines as selective VEGFR-2 and FGFR-1 kinase inhibitors for treatment of proliferative diseases
INVENTOR(S): Bhide, Rajeev; Ruel, Rejean; Thibeault, Carl; L'heureux, Alexandre
PATENT ASSIGNEE(S): Bristol-Myers Squibb Company, USA
SOURCE: PCT Int. Appl., 66 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 3
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2004009601	A1	20040129	WO 2003-US22554	20030718
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU				
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US 2004063707	A1	20040401	US 2003-622593	20030718
US 2004072832	A1	20040415	US 2003-623171	20030718

PRIORITY APPLN. INFO.: US 2002-397256P P 20020719
US 2003-447213P P 20030213

OTHER SOURCE(S): MARPAT 140:146173

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2004009601	A1	20040129	WO 2003-US22554	20030718
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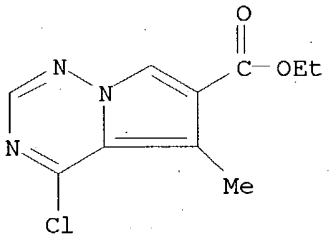
	RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC,				

NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG

US 2004063707 A1 20040401 US 2003-622593 20030718
US 2004072832 A1 20040415 US 2003-623171 20030718

IT **427878-41-9 649736-27-6 651744-49-9 651744-51-3**
RL: RCT (Reactant); RACT (Reactant or reagent)
(preparation of pyrrolotriazines as selective VEGFR-2 and FGFR-1 kinase inhibitors for treatment of proliferative diseases)

RN 427878-41-9 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 4-chloro-5-methyl-, ethyl ester (9CI) (CA INDEX NAME)



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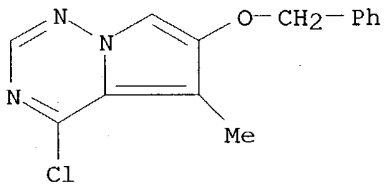
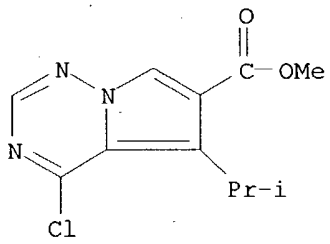
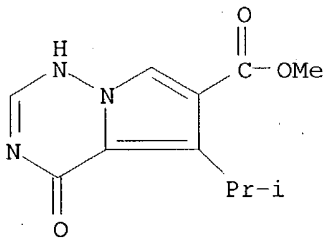
CN Pyrrolo[2,1-f][1,2,4]triazine, 4-chloro-5-methyl-6-(phenylmethoxy)- (9CI) (CA INDEX NAME)



RN 651744-49-9 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 4-chloro-5-(1-methylethyl)-, methyl ester (9CI) (CA INDEX NAME)

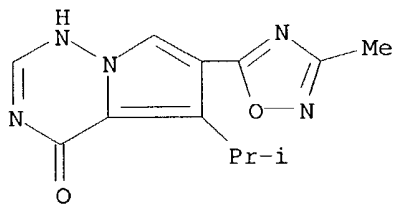


RN 651744-51-3 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-5-(1-methylethyl)-4-oxo-, methyl ester (9CI) (CA INDEX NAME)

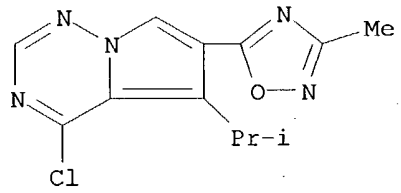


IT **651744-33-1P 651744-34-2P 651744-40-0P**
651753-52-5P 651753-54-7P
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(preparation of pyrrolotriazines as selective VEGFR-2 and FGFR-1 kinase inhibitors for treatment of proliferative diseases)

RN 651744-33-1 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazin-4(1H)-one, 5-(1-methylethyl)-6-(3-methyl-1,2,4-oxadiazol-5-yl)- (9CI) (CA INDEX NAME)



RN 651744-34-2 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine, 4-chloro-5-(1-methylethyl)-6-(3-methyl-1,2,4-oxadiazol-5-yl)- (9CI) (CA INDEX NAME)



RN 651744-40-0 CAPLUS
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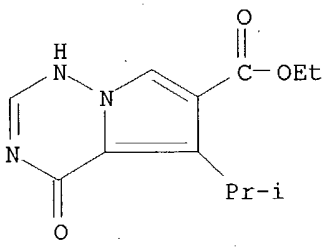
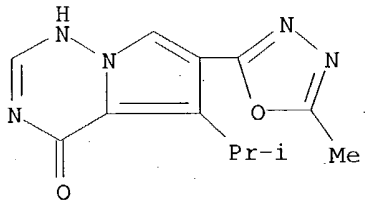
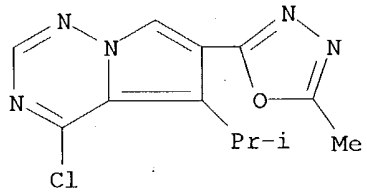


RN 651753-52-5 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazin-4(1H)-one, 5-(1-methylethyl)-6-(5-methyl-1,3,4-oxadiazol-2-yl)- (9CI) (CA INDEX NAME)



RN 651753-54-7 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine, 4-chloro-5-(1-methylethyl)-6-(5-methyl-1,3,4-oxadiazol-2-yl)- (9CI) (CA INDEX NAME)



REFERENCE COUNT: 1 THERE ARE 1 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L9 ANSWER 15 OF 15 CAPLUS COPYRIGHT 2004 ACS on STN
ACCESSION NUMBER: 2004:80644 CAPLUS
DOCUMENT NUMBER: 140:146018
TITLE: Process for preparation of indolyloxypyrrolotriazines and their use as drugs.
INVENTOR(S): Bhide, Rajeev; Fan, Junying; Parlanti, Luca; Barbosa, Stephanie; Qian, Ligang; Cai, Zhen-wei; Gibson, Francis S.
PATENT ASSIGNEE(S): Bristol-Myers Squibb Company, USA
SOURCE: PCT Int. Appl., 48 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 3
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2004009542	A2	20040129	WO 2003-US22755	20030721

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU

RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG

US 2004077858	A1	20040422	US 2003-622280	20030718

PRIORITY APPLN. INFO.:

US 2002-397256P	P	20020719
US 2003-447213P	P	20030213
US 2003-622280	A	20030718

OTHER SOURCE(S): MARPAT 140:146018

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2004009542	A2	20040129	WO 2003-US22755	20030721

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU

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US 2004077858	A1	20040422	US 2003-622280	20030718

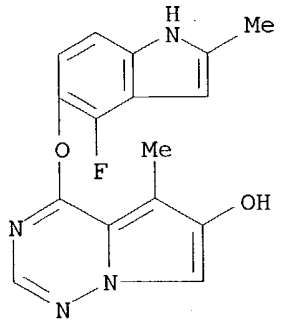
IT **649735-41-1P**

RL: IMF (Industrial manufacture); PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

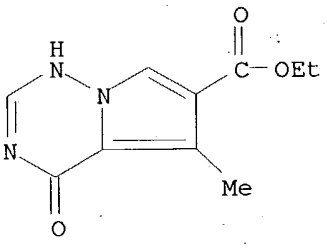
(process for preparation of indolyloxypyrrolotriazines and their use as drugs)

RN 649735-41-1 CAPLUS

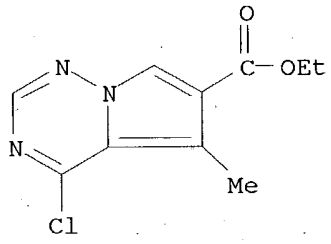
CN Pyrrolo[2,1-f][1,2,4]triazin-6-ol, 4-[(4-fluoro-2-methyl-1H-indol-5-yl)oxy]-5-methyl- (9CI) (CA INDEX NAME)



IT **427878-70-4**
RL: RCT (Reactant); RACT (Reactant or reagent)
(process for preparation of indolyloxypyrrolotriazines and their use as drugs)
RN 427878-70-4 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-5-methyl-4-oxo-, ethyl ester (9CI) (CA INDEX NAME)



IT **427878-41-9P 649736-26-5P 649736-27-6P**
649736-28-7P 649736-29-8P 649736-30-1P
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(process for preparation of indolyloxypyrrolotriazines and their use as drugs)
RN 427878-41-9 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 4-chloro-5-methyl-, ethyl ester (9CI) (CA INDEX NAME)



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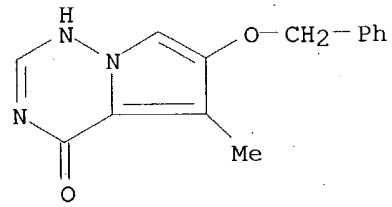

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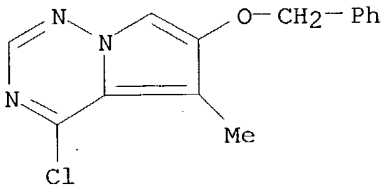
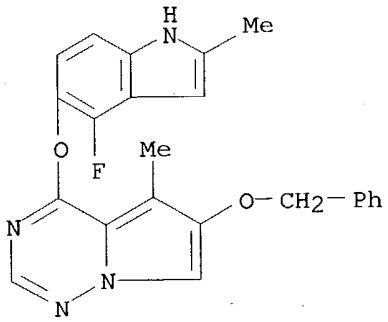
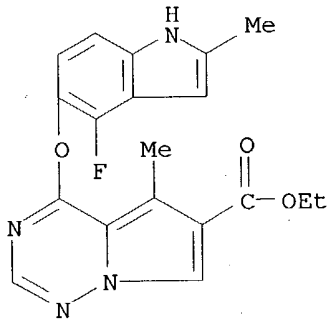


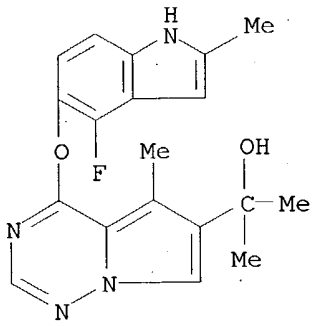
RN 649736-28-7 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine, 4-[(4-fluoro-2-methyl-1H-indol-5-yl)oxy]-5-methyl-6-(phenylmethoxy)- (9CI) (CA INDEX NAME)



RN 649736-29-8 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 4-[(4-fluoro-2-methyl-1H-indol-5-yl)oxy]-5-methyl-, ethyl ester (9CI) (CA INDEX NAME)



RN 649736-30-1 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-methanol, 4-[(4-fluoro-2-methyl-1H-indol-5-yl)oxy]-α,α,5-trimethyl- (9CI) (CA INDEX NAME)



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FILE COVERS 1907-1966
FILE LAST UPDATED:  01 May 1997 (19970501/UP)

  This file contains CAS Registry Numbers for easy and accurate
  substance identification.  Title keywords, authors, patent
  assignees, and patent information, e.g., patent numbers, are
  now searchable from 1907-1966.  TIFF images of CA abstracts
  printed between 1907-1966 are available in the PAGE
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  all substance data from the REGISTRY file.  Enter HELP FIRST for
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L10             0 L3

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LOGOFF? (Y)/N/HOLD:.
STN INTERNATIONAL LOGOFF AT 14:30:17 ON 11 MAY 2004
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10/62,3171 Thomas McKenzie

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NEWS 2		"Ask CAS" for self-help around the clock
NEWS 3	JAN 27	Source of Registration (SR) information in REGISTRY updated and searchable
NEWS 4	JAN 27	A new search aid, the Company Name Thesaurus, available in CA/CAplus
NEWS 5	FEB 05	German (DE) application and patent publication number format changes
NEWS 6	MAR 03	MEDLINE and LMEDLINE reloaded
NEWS 7	MAR 03	MEDLINE file segment of TOXCENTER reloaded
NEWS 8	MAR 03	FRANCEPAT now available on STN
NEWS 9	MAR 29	Pharmaceutical Substances (PS) now available on STN
NEWS 10	MAR 29	WPIFV now available on STN
NEWS 11	MAR 29	New monthly current-awareness alert (SDI) frequency in RAPRA
NEWS 12	APR 26	PROMT: New display field available
NEWS 13	APR 26	IFIPAT/IFIUDB/IFICDB: New super search and display field available
NEWS 14	APR 26	LITALERT now available on STN
NEWS 15	APR 27	NLDB: New search and display fields available
NEWS 16	May 10	PROUSDDR now available on STN
NEWS 17	May 10	PROUSDDR: One FREE connect hour, per account, in both May and June 2004

NEWS EXPRESS	MARCH 31 CURRENT WINDOWS VERSION IS V7.00A, CURRENT MACINTOSH VERSION IS V6.0c(ENG) AND V6.0Jc(JP), AND CURRENT DISCOVER FILE IS DATED 26 APRIL 2004
NEWS HOURS	STN Operating Hours Plus Help Desk Availability
NEWS INTER	General Internet Information
NEWS LOGIN	Welcome Banner and News Items
NEWS PHONE	Direct Dial and Telecommunication Network Access to STN
NEWS WWW	CAS World Wide Web Site (general information)

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FILE 'HOME' ENTERED AT 14:46:36 ON 11 MAY 2004

=> file reg
FILE 'REGISTRY' ENTERED AT 14:46:44 ON 11 MAY 2004
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STRUCTURE FILE UPDATES: 9 MAY 2004 HIGHEST RN 680971-82-8
DICTIONARY FILE UPDATES: 9 MAY 2004 HIGHEST RN 680971-82-8

TSCA INFORMATION NOW CURRENT THROUGH JANUARY 6, 2004

Please note that search-term pricing does apply when conducting SmartSELECT searches.

Crossover limits have been increased. See HELP CROSSOVER for details.

Experimental and calculated property data are now available. For more information enter HELP PROP at an arrow prompt in the file or refer to the file summary sheet on the web at:
http://www.cas.org/ONLINE/DBSS/registryss.html

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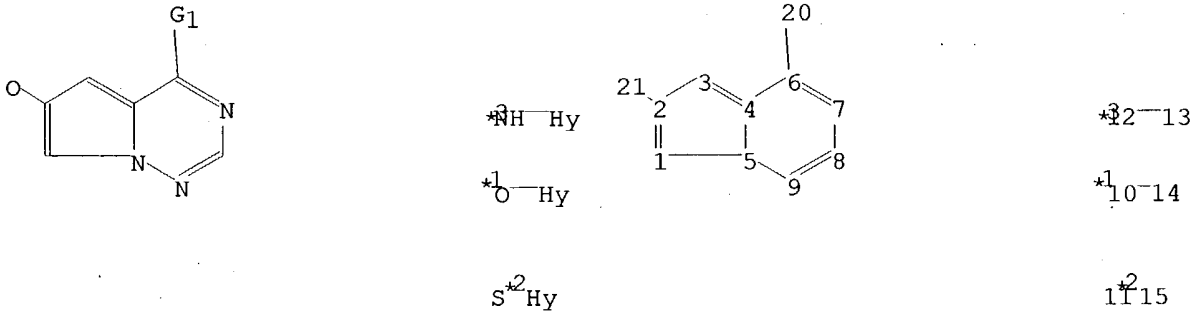

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ring nodes :
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ring bonds :
1-2 1-5 2-3 3-4 4-5 4-6 5-9 6-7 7-8 8-9
exact/norm bonds :
1-2 1-5 2-3 2-21 3-4 4-5 4-6 5-9 6-7 6-20 7-8 8-9 10-14 11-15 12-13

G1:OH,Cl,[*1],[*2],[*3]

Match level :
1:Atom 2:Atom 3:Atom 4:Atom 5:Atom 6:Atom 7:Atom 8:Atom 9:Atom 10:CLASS
11:CLASS 12:CLASS 13:Atom 14:Atom 15:Atom 20:CLASS 21:CLASS
Generic attributes :
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Number of Carbon Atoms : 7 or more
Number of Hetero Atoms : less than 2
Type of Ring System : Polycyclic
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Saturation : Unsaturated
Number of Carbon Atoms : 7 or more
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Type of Ring System : Polycyclic
15:
Saturation : Unsaturated
Number of Carbon Atoms : 7 or more
Number of Hetero Atoms : less than 2
Type of Ring System : Polycyclic

L1 STRUCTURE UPLOADED

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FULL SCREEN SEARCH COMPLETED - 310 TO ITERATE

100.0% PROCESSED 310 ITERATIONS 96 ANSWERS
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=> file caplus
FILE 'CAPLUS' ENTERED AT 14:47:36 ON 11 MAY 2004
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FILE COVERS 1907 - 11 May 2004 VOL 140 ISS 20
FILE LAST UPDATED: 10 May 2004 (20040510/ED)

This file contains CAS Registry Numbers for easy and accurate
substance identification.

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L4   ANSWER 1 OF 4  CAPLUS  COPYRIGHT 2004 ACS on STN
ACCESSION NUMBER:         2004:120859  CAPLUS
DOCUMENT NUMBER:          140:181471
TITLE:                    Preparation of pyrrolotriazines as tyrosine kinase
                          activity inhibitors of growth factor receptors for the
                          treatment of cancer
INVENTOR(S):              Bhide, Rajeev S.; Borzilleri, Robert M.
PATENT ASSIGNEE(S):       Bristol-Myers Squibb Company, USA
SOURCE:                   PCT Int. Appl., 71 pp.
                          CODEN: PIXXD2
DOCUMENT TYPE:            Patent
LANGUAGE:                 English
FAMILY ACC. NUM. COUNT:   1
PATENT INFORMATION:

     PATENT NO.        KIND  DATE           APPLICATION NO.   DATE
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     WO 2004013145      A1   20040212       WO 2003-US24273   20030804
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             CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH,
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             LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM,
             PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN,
             TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW, AM, AZ, BY,
             KG, KZ, MD, RU
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             CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC,
             NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ,
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     US 2004063708      A1   20040401       US 2003-633997    20030804
PRIORITY APPLN. INFO.:                   US 2002-400572P  P  20020802
OTHER SOURCE(S):          MARPAT 140:181471
     PATENT NO.        KIND  DATE           APPLICATION NO.   DATE
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PI   WO 2004013145      A1   20040212       WO 2003-US24273   20030804
         W:  AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN,
             CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH,
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             TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW, AM, AZ, BY,
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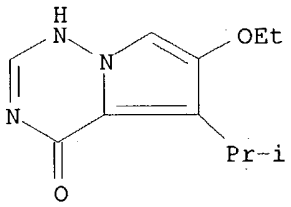
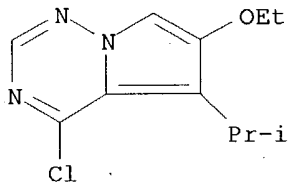
NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG
US 2004063708 A1 20040401 US 2003-633997 20030804

IT **658085-59-7P 658085-60-0P**
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(intermediate; preparation of pyrrolotriazines as tyrosine kinase activity inhibitors of growth factor receptors)

RN 658085-59-7 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazin-4(1H)-one, 6-ethoxy-5-(1-methylethyl)- (9CI) (CA INDEX NAME)



RN 658085-60-0 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine, 4-chloro-6-ethoxy-5-(1-methylethyl)- (9CI) (CA INDEX NAME)



L4 ANSWER 2 OF 4 CAPLUS COPYRIGHT 2004 ACS on STN
ACCESSION NUMBER: 2004:80698 CAPLUS
DOCUMENT NUMBER: 140:146173
TITLE: Preparation of pyrrolotriazines as selective VEGFR-2 and FGFR-1 kinase inhibitors for treatment of proliferative diseases
INVENTOR(S): Bhide, Rajeev; Ruel, Rejean; Thibeault, Carl; L'heureux, Alexandre
PATENT ASSIGNEE(S): Bristol-Myers Squibb Company, USA
SOURCE: PCT Int. Appl., 66 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 3
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2004009601	A1	20040129	WO 2003-US22554	20030718

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GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR,
LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM,
PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN,
TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW, AM, AZ, BY,
KG, KZ, MD, RU
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG,
CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC,
NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ,
GW, ML, MR, NE, SN, TD, TG

US 2004063707	A1	20040401	US 2003-622593	20030718	
US 2004072832	A1	20040415	US 2003-623171	20030718	

PRIORITY APPLN. INFO.: US 2002-397256P P 20020719
US 2003-447213P P 20030213

OTHER SOURCE(S): MARPAT 140:146173

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI WO 2004009601	A1	20040129	WO 2003-US22554	20030718

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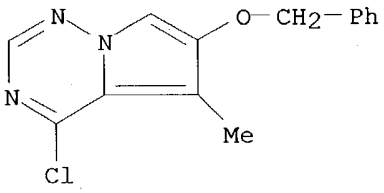
CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC,
NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ,
GW, ML, MR, NE, SN, TD, TG

US 2004063707	A1	20040401	US 2003-622593	20030718	
US 2004072832	A1	20040415	US 2003-623171	20030718	

IT **649736-27-6**
RL: RCT (Reactant); RACT (Reactant or reagent)
(preparation of pyrrolotriazines as selective VEGFR-2 and FGFR-1 kinase inhibitors for treatment of proliferative diseases)

RN 649736-27-6 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine, 4-chloro-5-methyl-6-(phenylmethoxy)- (9CI) (CA INDEX NAME)



REFERENCE COUNT: 1 THERE ARE 1 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 3 OF 4 CAPLUS COPYRIGHT 2004 ACS on STN
ACCESSION NUMBER: 2004:80644 CAPLUS
DOCUMENT NUMBER: 140:146018
TITLE: Process for preparation of indolyloxypyrrolotriazines and their use as drugs.
INVENTOR(S): Bhide, Rajeev; Fan, Junying; Parlanti, Luca; Barbosa, Stephanie; Qian, Ligang; Cai, Zhen-wei; Gibson,

Francis S.
PATENT ASSIGNEE(S): Bristol-Myers Squibb Company, USA
SOURCE: PCT Int. Appl., 48 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 3
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2004009542	A2	20040129	WO 2003-US22755	20030721

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU

RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG

US 2004077858	A1	20040422	US 2003-622280	20030718

PRIORITY APPLN. INFO.:
US 2002-397256P P 20020719
US 2003-447213P P 20030213
US 2003-622280 A 20030718

OTHER SOURCE(S): MARPAT 140:146018

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2004009542	A2	20040129	WO 2003-US22755	20030721

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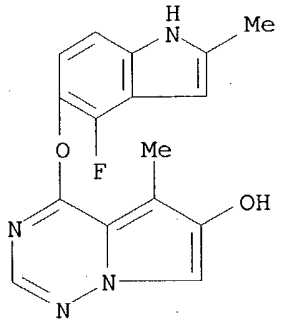
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG

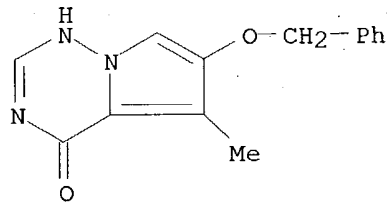
US 2004077858	A1	20040422	US 2003-622280	20030718

IT **649735-41-1P**
RL: IMF (Industrial manufacture); PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(process for preparation of indolyloxypyrrolotriazines and their use as drugs)

RN 649735-41-1 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazin-6-ol, 4-[(4-fluoro-2-methyl-1H-indol-5-yl)oxy]-5-methyl- (9CI) (CA INDEX NAME)



IT **649736-26-5P 649736-27-6P 649736-28-7P**
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(process for preparation of indolyloxypyrrolotriazines and their use as drugs)
RN 649736-26-5 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazin-4(1H)-one, 5-methyl-6-(phenylmethoxy)- (9CI) (CA INDEX NAME)



RN 649736-27-6 CAPLUS
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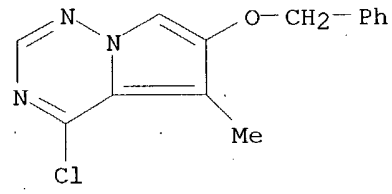
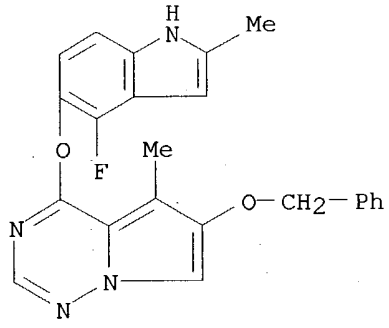


RN 649736-28-7 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine, 4-[(4-fluoro-2-methyl-1H-indol-5-yl)oxy]-5-methyl-6-(phenylmethoxy)- (9CI) (CA INDEX NAME)



L4 ANSWER 4 OF 4 CAPLUS COPYRIGHT 2004 ACS on STN
ACCESSION NUMBER: 2002:391720 CAPLUS
DOCUMENT NUMBER: 136:386144
TITLE: Preparation of pyrrolo[2,1-f][1,2,4]triazine carboxylic acid derivatives for use in treating p38 kinase-associated conditions
INVENTOR(S): Leftheris, Katerina; Barrish, Joel; Hynes, John; Wrobleski, Stephen T.
PATENT ASSIGNEE(S): Bristol-Myers Squibb Company, USA
SOURCE: PCT Int. Appl., 108 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 2
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2002040486	A2	20020523	WO 2001-US49982	20011107
WO 2002040486	A3	20030912		
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RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
AU 2002032760	A5	20020527	AU 2002-32760	20011107
EE 200300227	A	20031015	EE 2003-227	20011107
EP 1363910	A2	20031126	EP 2001-992298	20011107
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NO 2003002229	A	20030716	NO 2003-2229	20030516

PRIORITY APPLN. INFO.:
US 2000-249877P P 20001117
US 2001-310561P P 20010807
WO 2001-US49982 W 20011107

OTHER SOURCE(S): MARPAT 136:386144

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2002040486	A2	20020523	WO 2001-US49982	20011107
	WO 2002040486	A3	20030912		

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AU 2002032760	A5	20020527	AU 2002-32760	20011107	
EE 200300227	A	20031015	EE 2003-227	20011107	
EP 1363910	A2	20031126	EP 2001-992298	20011107	
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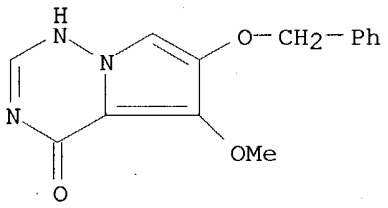
NO 2003002229	A	20030716	NO 2003-2229	20030516	

IT **310444-95-2P 310444-96-3P**, 4-Chloro-5-methoxy-6-[phenylmethoxy]pyrrolo[2,1-f][1,2,4]triazine
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(intermediate; preparation of pyrrolo[2,1-f][1,2,4]triazine carboxylic acid derivs. for use in treating p38 kinase-associated conditions)

RN 310444-95-2 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazin-4(1H)-one, 5-methoxy-6-(phenylmethoxy)- (9CI) (CA INDEX NAME)



RN 310444-96-3 CAPLUS
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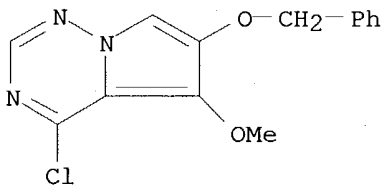

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